(FILE 'HOME' ENTERED AT 14:45:14 ON 11 MAY 2003) FILE 'REGISTRY' ENTERED AT 14:45:19 ON 11 MAY 2003 5 S IPA/CN L11 s (ISOPROPANOL)/CN L2 FILE 'CAPLUS' ENTERED AT 14:45:58 ON 11 MAY 2003 42546 S L2 L3 L4 1203 S L3 AND (?SILOX?) 0 S (CERAMIC HYDBRID?) L5 109 S (CERAMIC HYBRID?) Lέ L7 0 S L4 AND L6 0 S L4 AND (CIRCUIT? AND CERAMIC?) L8 16 S L4 AND CIRCUIT? L9 16 FOCUS L9 1-L1016 S L4 AND HYBRID? L1115 S L11 NOT L9 L12 15 FOCUS L12 1-L13

=>

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L10 ANSWER 1 OF 16 CAPLUS COPYRIGHT 2003 ACS
    1989:9818 CAPLUS
AN
    110:9818
DN
    Storage-stable ladder siloxane coating materials
ΤI
    Nagata, Kenji
ΙN
    Fujitsu Ltd., Japan
PA
    Jpn. Kokai Tokkyo Koho, 4 pp.
SO
    CODEN: JKXXAF
DT
    Patent
    Japanese
LA
    ICM C09D003-82
ΙC
ICA C08G077-38
     42-10 (Coatings, Inks, and Related Products)
     Section cross-reference(s): 76
FAN.CNT 1
                                         APPLICATION NO. DATE
     PATENT NO.
                    KIND DATE
                                          _____
     ______
                                         JP 1987-30055
                                                           19870212
PΤ
    JP 63196669
                     A2 19880815
                           19870212
PRAI JP 1987-30055
    Title materials, useful as insulation films in multilayer printed
     circuits, are prepd. by self condensation of poly(
     diethoxysiloxane) (I) during solvent exchange from isopropanol to
     butyl Cellosolve acetate (II) to decrease OH content in I, then treating
     the resulting polymer with glass resins. Thus, a soln. of 300 .+-. 1.5~\mathrm{g}
     I in isopropanol was mixed with 1635 .+-. 0.8 g II and vacuum evapd. at 35
     .+-. 1.degree., then the resulting mixt. was treated with 48.0 .+-. 0.2 g
     glass resin at 60.degree. to give a ladder siloxane, which
     showed no change in viscosity when stored at 5.degree. for 1 mo.
     ladder siloxane insulator printed circuit; glass resin
ST
     coating printed circuit; printed circuit insulating
     coating siloxane
ΙT
     Electric insulators and Dielectrics
        (coatings, ladder siloxanes, prepn. of, with good storage
        stability, for printed circuits)
ΙT
     Electric circuits
        (printed, insulator films for, ladder siloxanes as, with good
        storage stability)
                                                  112-07-2, Butyl
     67-63-0, Isopropanol, uses and miscellaneous
ΙT
     cellosolve acetate
     RL: USES (Uses)
        (solvent, in manuf. of storage-stable ladder siloxane
        coatings)
RN
     67-63-0
     112-07-2
RN
L10 ANSWER 2 OF 16 CAPLUS COPYRIGHT 2003 ACS
     1996:472864 CAPLUS
AN
     125:171569
DN
     Environment-friendly washing process using siloxanes
ТŢ
     Shirai, Michio
ΤN
     Olympus Optical Co, Japan
PΑ
SO
     Jpn. Kokai Tokkyo Koho, 10 pp.
     CODEN: JKXXAF
DT
     Patent
LA
     Japanese
     ICM C11D007-22
TC
     TCS B08B003-08
     46-6 (Surface Active Agents and Detergents)
     Section cross-reference(s): 48, 76
FAN.CNT 1
```

```
PATENT NO. KIND DATE
                                          APPLICATION NO. DATE
                                           _____
     _____
                      ____
     JP 08134498
                    A2 19960528
19941110
                                          JP 1994-276603 19941110
PRAI JP 1994-276603
     For diagram(s), see printed CA Issue.
     Washing liqs. contg. low-mol. volatile Me3SiO(SiMe2O)nSiMe3 (n = 0, 1, 2)
AΒ
     and/or cyclic siloxanes I (m = 4, 5) in consecutively linked
     washing baths are moved reversely toward the moving direction of the
     substrates, and the liqs. are removed from the 1st bath, distd. to remove
     highly volatile impurities, and supplied to the last bath. The washing
     liqs. may be (pseudo)azeotropic (1) a mixt. of
     hexamethyldisiloxane and MeOH, EtOH, or isopropanol, (2) a mixt.
     of octamethyltrisiloxane (II) and isopentyl alc., 2-pentanol,
     propylene glycol mono-Et ether, propylene glycol tert-Bu ether, propylene
     glycol mono-Me ether acetate (III), propylene glycol mono-Et ether acetate
     (IV), or 3-methyl-3-methoxybutyl acetate, (3) a mixt. of
     decamethyltetrasiloxane and dihydroterpineol, or (4) a mixt. of
     octamethylcyclotetrasiloxane and III or IV. Lead frames of Cu
     were subjected to the washing process using II alone.
     environment friendly washing process siloxane; lead frame copper
     washing process; cyclic siloxane detergent; azeotropic
     siloxane alc mixt detergent; pseudoazeotropic siloxane
     alc mixt detergent
ΙT
     Detergents
     Electric circuits
        (low-mol.-wt. volatile (cyclic) siloxanes or their
        (pseudo)azeotropic mixts. with alcs. as recyclable washing ligs.)
ΙT
     Alcohols, uses
       Cyclosiloxanes
       Siloxanes and Silicones, uses
     RL: PEP (Physical, engineering or chemical process); TEM (Technical or
     engineered material use); PROC (Process); USES (Uses)
        (low-mol.-wt. volatile (cyclic) siloxanes or their
        (pseudo)azeotropic mixts. with alcs. as recyclable washing liqs.)
     64-17-5, Ethanol, uses 67-56-1, Methanol, uses 67-63-0,
ΙT
     Isopropanol, uses 107-46-0, Hexamethyldisiloxane 107-51-7, Octamethyltrisiloxane 123-51-3, Isopentyl alcohol 141-62-8,
     Decamethyltetrasiloxane 541-02-6,
     Decamethylcyclopentasiloxane 556-67-2,
Octamethylcyclotetrasiloxane 6032-29-7, 2-Pentanol 52125-53-8,
     Propylene glycol monoethyl ether 58985-02-7, Dihydroterpineol
     80763-10-6 84540-57-8, Propylene glycol monomethyl ether acetate
     98516-30-4, Propylene glycol monoethyl ether acetate 103429-90-9,
     3-Methyl-3-methoxybutyl acetate
     RL: PEP (Physical, engineering or chemical process); TEM (Technical or
     engineered material use); PROC (Process); USES (Uses)
        (low-mol.-wt. volatile (cyclic) siloxanes or their
        (pseudo)azeotropic mixts. with alcs. as recyclable washing liqs.)
RN
     64-17-5
RN
    67-56-1
RN
    67-63-0
RN
    107-46-0
RN
    107-51-7
RN
    123-51-3
RN
   141-62-8
RN
   541-02-6
RN 556-67-2
RN 6032-29-7
RN 52125-53-8
RN 58985-02-7
RN 80763-10-6
RN 84540-57-8
```

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RN 103429-90-9
L10 ANSWER 3 OF 16 CAPLUS COPYRIGHT 2003 ACS
    1986:544619 CAPLUS
DN
    105:144619
    Cleaning a circuit board
TΙ
    Wong, Ching Ping
IN
PA
   AT and T Technologies, Inc., USA
    U.S., 3 pp.
SC
    CODEN: USXXAM
DT
    Patent
    English
LA
ΙC
    ICM C23G001-02
    ICS C23G001-14
NCL 134028000
    76-14 (Electric Phenomena)
CC
FAN.CNT 1
                                          APPLICATION NO. DATE
    PATENT NO.
                    KIND DATE
     _____
                                          _____
                                          US 1985-774674 19850911
    US 4604144
                    A 19860805
                          19850911
PRAI US 1985-774674
    In a method of cleaning a circuit board having a
     silicone-encapsulated hybrid integrated circuit on it,
     subsequent to aq.-solder-flux residue removal with an ionic detergent
     (e.g., ethanolamine) the board is cleaned with a low-mol.-wt. org. acid,
     such as formic, acetic or oxalic acid. Addnl., the silicone may be
     swelled prior to or during treatment with the aq. org. acid soln.
     swellant comprises a member of the group consisting of alcs. and
     fluorocarbons, including MeOH, EtOH, PrOH and i-PrOH. Application of the
    method to cleaning hybrid integrated circuits is indicated.
     circuit board cleaning; carboxylic acid circuit board
ST
     cleaning; silicone encapsulant circuit board cleaning; ionic
     detergent circuit board cleaning; integrated circuit
     cleaning
ΙT
     Soldering
        (elec. circuit board cleaning prior to)
     Siloxanes and Silicones, uses and miscellaneous
ΙT
     RL: USES (Uses)
        (encapsulants, for elec. circuit boards, cleaning of)
     Carboxylic acids, uses and miscellaneous
ΙT
     RL: USES (Uses)
        (in elec. circuit board cleaning)
ΙT
     Alcohols, uses and miscellaneous
     RL: USES (Uses)
        (swellants, for silicone encapsulants, in elec. circuit board
        cleaning)
     Hydrocarbons, uses and miscellaneous
ΙT
     RL: USES (Uses)
        (chloro fluoro, swellants, for silicon encapsulants, in elec.
        circuit board cleaning)
TT
     Electric circuits
        (integrated, cleaning of)
TT
     Detergents
        (ionic, in elec. circuit board cleaning)
     Electric circuits
IT
        (printed, boards, cleaning of)
     141-43-5, uses and miscellaneous
TΤ
     RL: USES (Uses)
        (detergent, for elec. circuit board cleaning)
                                     64-19-7, uses and miscellaneous
     64-18-6, uses and miscellaneous
ΙT
     144-62-7, uses and miscellaneous
```

RN

98516-30-4

```
RL: USES (Uses)
       (in elec. circuit board cleaning)
    64-17-5, uses and miscellaneous
                                      67-56-1, uses and miscellaneous
ΙT
     67-63-0, uses and miscellaneous
                                      71-23-8, uses and miscellaneous
    RL: USES (Uses)
       (swellant, for silicone encapsulants, in elec. circuit board
       cleaning)
    141-43-5
RN
RN
    64-18-6
    64-19-7
RN
    144-62-7
RN
    64-17-5
RN
    67-56-1
RN
    67-63-0
RN
    71-23-8
RN
L10 ANSWER 4 OF 16 CAPLUS COPYRIGHT 2003 ACS
     1983:99846 CAPLUS
AN
     98:99846
DN
     Printed circuit board
TΙ
    Fujikura Cable Works, Ltd., Japan
PΑ
     Jpn. Kokai Tokkyo Koho, 6 pp.
SO
     CODEN: JKXXAF
DT
     Patent
     Japanese
A.T
     H05K003-44
TC
     76-14 (Electric Phenomena)
CC
FAN.CNT 1
                                           APPLICATION NO.
                                                            DATE
                     KIND DATE
     PATENT NO.
                           _____
     _____
                                           JP 1981-56816
                                                            19810415
                            19821023
     JP 57172797
                      A2
PΤ
PRAI JP 1981-56816
                            19810415
     The printed circuit board is characterized by a <0.4 mm thick Al
     layer formed on the surface of the metal such as Fe, Fe alloy, Cu, Cu
     alloy or Al-Si based alloy, and anodically formed pores filled with a
     polymer.
     alumina polymer insulator printed circuit
ST
     Siloxanes and Silicones, uses and miscellaneous
IT
     RL: TEM (Technical or engineered material use); USES (Uses)
        (elec. insulators, for printed-circuit board)
     Electric circuits
ΙT
        (printed, boards, alumina and polymer insulators for)
     2171-98-4D, hyd., polymer 6184-20-9D, hyd., polymer 51382-55-9D, hyd.,
ΙT
             83644-71-7D, hyd., polymer
     RL: TEM (Technical or engineered material use); USES (Uses)
        (elec. insulators, for printed-circuit board)
     64-19-7, uses and miscellaneous 67-63-0, uses and miscellaneous
IT
     RL: USES (Uses)
        (in prepn. of porous substrates for printed circuits)
     7429-90-5, uses and miscellaneous 7439-89-6, uses and miscellaneous
ΤТ
     7440-50-8, uses and miscellaneous
     RL: USES (Uses)
        (on aluminum alloy porous boards for printed circuits)
     12679-19-5
IT
     RL: USES (Uses)
        (on porous board for printed circuits)
     2171-98-4D
RN
RN
     6184-20-9D
     51382-55-9D
RN
RN
     83644-71-7D
RN
    64-19-7
RN
     67-63-0
```

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7439-89-6
RN
     7440-50-8
RN
    12679-19-5
RN
L10 ANSWER 5 OF 16 CAPLUS COPYRIGHT 2003 ACS
     1993:130098 CAPLUS
ΑN
     118:130098
DΝ
     Silicone resin additives for fluxes and soldering pastes
TΙ
     Gomi, Tadashi; Douzaki, Yuji
IN
     Yuho Chemicals Inc., Japan
PΑ
     Eur. Pat. Appl., 13 pp.
SO
     CODEN: EPXXDW
DT
     Patent
     English
LA
     ICM B23K035-363
ΙC
     56-9 (Nonferrous Metals and Alloys)
     Section cross-reference(s): 76
FAN.CNT 1
                                          APPLICATION NO. DATE
                     KIND DATE
     PATENT NO.
                           _____
     _____ ___
                                           _____
                                                           19920416
                                           EP 1992-106665
                      A1
                            19921028
     EP 510539
PΤ
     EP 510539
                     В1
                            19970827
        R: DE, FR, GB, NL, SE
                                           JP 1991-112550
                                                           19910417
     JP 05092296 A2
                            19930416
                            19960124
     JP 08004953
                      B4
                                           US 1992-868416 19920415
     US 5215601
                      Α
                            19930601
PRAI JP 1991-112550
                            19910417
     The soldering flux mixts. and pastes include silicone resins to react with
     org. acids or their salts for deactivation and decreased corrosion. The
     resins are selected from methylsilicone, methylphenylsilicone, and/or
     modified silicone types. The flux mixt. optionally includes a solvent
     with s.p. below the m.p. of solder. The modified flux preferably has the
     COOH: SiOH mol ratio of 1: (0.5-1.5), and is suitable for the pastes contg.
     40-95 wt.* solder. The preferred fluxes contain org. acids (including
     rosins) and their salts 0.1-20, silicone resin 0.05-80, and solvents
     10-99.85 \text{ wt.-}. The solder pastes show decreased corrosion of Cu, esp. for
     elec.-circuit boards.
     solder paste flux silicone resin; org acid flux deactivation silicone;
ST
     rosin flux deactivation silicone; elec circuit soldering flux
     silicone
     Carboxylic acids, uses
IT
     Rosin
     RL: USES (Uses)
        (flux mixts. contg., for solder pastes, org. acids neutralized by
        silicone resin in)
     Siloxanes and Silicones, uses
ΙT
     RL: USES (Uses)
         (flux mixts. contg., to neutralize org. acids in soldering)
     Solders
IΤ
        (pastes, silicone resins in fluxes for, to neutralize org. acids and
        rosins)
     Siloxanes and Silicones, uses
TT
     RL: USES (Uses)
        (alkyd-, flux mixts. contg., for solder pastes, org. acids neutralized
        by silicone resin in)
     Siloxanes and Silicones, uses
IΤ
     RL: USES (Uses)
        (epoxy, flux mixts. contg., for solder pastes, org. acids neutralized
        by silicone resin in)
     Soldering
ΙT
         (fluxes, silicone resins in, to neutralize org. acids and rosins)
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7429-90-5

RN

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Castor oil
ΙT
    RL: USES (Uses)
        (hydrogenated, flux mixts. contg., for solder pastes, org. acids
        neutralized by silicone resin in)
    Siloxanes and Silicones, uses
ΙT
    RL: USES (Uses)
        (hydroxy-contg., flux mixts. contg., for solder pastes, org. acids
        neutralized by silicone resin in)
    Siloxanes and Silicones, uses
TT
    RL: USES (Uses)
        (methoxy, flux mixts. contg., for solder pastes, org. acids neutralized
        by silicone resin in)
IT
     Electric circuits
        (printed, boards, soldering of, fluxes and pastes for, with silicone
        resins)
    Alkyd resins
IΤ
     Epoxy resins, uses
     RL: USES (Uses)
        (siloxane-, flux mixts. contg., for solder pastes, org. acids
        neutralized by silicone resin in)
     64-17-5, Ethanol, uses 67-63-0, Isopropanol, uses 79-14-1,
     Hydroxyacetic acid, uses 110-15-6, Succinic acid, uses 110-94-1,
     Glutaric acid 111-20-6, Sebacic acid, uses 123-99-9, Azelaic acid,
           124-04-9, Adipic acid, uses 141-82-2, Malonic acid, uses
                                                                     137592-56-4
     34590-94-8, Dipropylene glycol monomethyl ether
                                                       84376-09-0
     146448-14-8
     RL: USES (Uses)
        (flux mixts. contg., for solder pastes, org. acids neutralized by
        silicone resin in)
     514-10-3, Abietic acid
IT
     RL: USES (Uses)
        (flux mixts. contg., for soldering, silicone resins for acidity
        neutralization in)
                               136766-80-8
                                            136766-83-1
                 62258-61-1
     12610-63-8
ΙT
     RL: USES (Uses)
        (solder, flux mixt. for pastes contg. powd., with silicone resin to
        neutralize org. acids)
ΙT
     7440-50-8, Copper, uses
     RL: USES (Uses)
        (soldering of, flux mixts. in, with silicone resins to neutralize org.
        acids and rosin)
     64-17-5
RN
     67-63-0
RN
     79-14-1
RN
     110-15-6
RN
RN
     110-94-1
RN
     111-20-6
RN
     123-99-9
RN
     124-04-9
     141-82-2
RN
     34590-94-8
RN
     84376-09-0
RN
RN
     137592-56-4
RN
     146448-14-8
RN
     514-10-3
     12610-63-8
RN
     62258-61-1
RN
     136766-80-8
RN
     136766-83-1
RN
     7440-50-8
RN
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L10 ANSWER 6 OF 16 CAPLUS COPYRIGHT 2003 ACS

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2002:669624 CAPLUS
AN
DN
    137:194031
    Dielectric films for narrow gap-fill applications in integrated-
TΙ
     circuit fabrication
     Leung, Roger; Endisch, Denis; Xie, Songyuan; Hacker, Nigel; Deng, Yanpei
IN
     Honeywell International, Inc., USA
    U.S., 11 pp.
SO
     CODEN: USXXAM
\mathsf{D}\mathbf{T}
    Patent
     English
LA
     ICM H01L021-44
T.C.
NCL 438118000
     76-3 (Electric Phenomena)
CC
FAN.CNT 1
                                         APPLICATION NO. DATE
                     KIND DATE
     PATENT NO.
                                           ______
                     ____
                                          US 2001-761529
                                                            20010111
                     B1 20020903
     US 6444495
PΤ
     US 2002137260
                     A1 20020926
                                          US 2002-188433 20020703
                     A1 20030508
     US 2003087485
PRAI US 2001-761529 A3
                          20010111
    A colloidal suspension of nanoparticles composed of a dense material
     dispersed in a solvent was used in forming a gap-filling dielec. material
     with low thermal shrinkage, high thermal stability, and high etching
     resistance. The dielec. material is particularly useful for pre-metal
     dielec. and shallow trench isolation applications. According to the
     methods of forming a dielec. material, the colloidal suspension is
     deposited on a substrate and dried to form a porous intermediate layer.
     The intermediate layer is modified by infiltration with a liq. phase
     matrix material, such as a spin-on polymer, followed by curing, by
     infiltration with a gas phase matrix material, followed by curing, or by
     curing alone, to provide a gap-filling, thermally stable, etch resistant
     dielec. material.
     colloidal dielec film spin coating integrated circuit
ST
     fabrication
     Silsesquioxanes
     RL: PEP (Physical, engineering or chemical process); PYP (Physical
     process); SPN (Synthetic preparation); TEM (Technical or engineered
     material use); PREP (Preparation); PROC (Process); USES (Uses)
        (boron-doped; colloidal dielec. films for narrow gap-fill applications
        in integrated-circuit fabrication)
     Annealing
TΤ
     Colloids
     Dielectric films
     Drying
     Integrated circuits
     Nitriding
     Oxidation
        (colloidal dielec. films for narrow gap-fill applications in
        integrated-circuit fabrication)
     Polymers, processes
     RL: PEP (Physical, engineering or chemical process); PYP (Physical
     process); TEM (Technical or engineered material use); PROC (Process); USES
     (Uses)
        (colloidal dielec. films for narrow gap-fill applications in
        integrated-circuit fabrication)
     Borosilicates
TT
     RL: SPN (Synthetic preparation); TEM (Technical or engineered material
     use); PREP (Preparation); USES (Uses)
        (colloidal dielec. films for narrow gap-fill applications in
        integrated-circuit fabrication)
     Borophosphosilicate glasses
TT
     Borosilicate glasses
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Polycarbosilanes
       Polysiloxanes, uses
     Silazanes
     RL: TEM (Technical or engineered material use); USES (Uses)
        (colloidal dielec. films for narrow gap-fill applications in
        integrated-circuit fabrication)
ΙT
     Porous materials
        (films; colloidal dielec. films for narrow gap-fill applications in
        integrated-circuit fabrication)
TΤ
     Silicates, uses
     RL: SPN (Synthetic preparation); TEM (Technical or engineered material
     use); PREP (Preparation); USES (Uses)
        (phospho-; colloidal dielec. films for narrow gap-fill applications in
        integrated-circuit fabrication)
ΙT
     Films
        (porous; colloidal dielec. films for narrow gap-fill applications in
        integrated-circuit fabrication)
ΤТ
     Coating process
        (spin; colloidal dielec. films for narrow gap-fill applications in
        integrated-circuit fabrication)
TΤ
     998-30-1, Triethoxysilane
     RL: CPS (Chemical process); NUU (Other use, unclassified); PEP (Physical,
     engineering or chemical process); PROC (Process); USES (Uses)
        (colloidal dielec. films for narrow gap-fill applications in
        integrated-circuit fabrication)
     7440-36-0, Antimony, uses 7440-38-2, Arsenic, uses 7440-42-8, Boron,
ΙT
           7723-14-0, Phosphorus, uses
     RL: MOA (Modifier or additive use); USES (Uses)
        (colloidal dielec. films for narrow gap-fill applications in
        integrated-circuit fabrication)
     64-17-5, Ethanol, uses 67-63-0, Isopropyl alcohol, uses
IT
                                                              1303-86-2, Boron
     67-64-1, Acetone, uses 108-94-1, Cyclohexanone, uses
     oxide (B2O3), uses 7697-37-2, Nitric acid, uses 7727-37-9, Nitrogen,
            30136-13-1, Propoxypropanol
     RL: NUU (Other use, unclassified); USES (Uses)
        (colloidal dielec. films for narrow gap-fill applications in
        integrated-circuit fabrication)
                                   12125-01-8P, Ammonium fluoride
     7631-86-9P, Silica, processes
     RL: PEP (Physical, engineering or chemical process); PYP (Physical
     process); SPN (Synthetic preparation); TEM (Technical or engineered
     material use); PREP (Preparation); PROC (Process); USES (Uses)
        (colloidal dielec. films for narrow gap-fill applications in
        integrated-circuit fabrication)
                                                                 12033-89-5P,
                                11105-01-4P, Silicon oxynitride
TΤ
     7440-21-3P, Silicon, uses
     Silicon nitride, uses
     RL: SPN (Synthetic preparation); TEM (Technical or engineered material
     use); PREP (Preparation); USES (Uses)
        (colloidal dielec. films for narrow gap-fill applications in
        integrated-circuit fabrication)
     1344-28-1, Aluminum oxide, uses 6876-41-1D, derivs.
                                                            7429-90-5,
ŢΤ
                      24304-00-5, Aluminum nitride 153315-81-2, Hydrogen
     Aluminum, uses
     silsesquioxane
     RL: TEM (Technical or engineered material use); USES (Uses)
        (colloidal dielec. films for narrow gap-fill applications in
        integrated-circuit fabrication)
              THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT 10
RE.
(1) Fischer; US 6143401 A 2000
(2) Jeng; US 5548159 A 1996
(3) Jeng; US 5858871 A 1999
(4) Mushiake; US 6242135 Bl 2001 CAPLUS
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Phosphosilicate glasses

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(5) Rosenmayer; US 5889104 A 1999 CAPLUS
(6) Ryang; US 5962608 A 1999 CAPLUS
(7) Smith; US 6319852 B1 2001 CAPLUS
(8) Stober; US 3634558 A 1972 CAPLUS
(9) Yamada; US 6245439 B1 2001
(10) Zakhidov; US 6261469 B1 2001 CAPLUS
    998-30-1
RN
    7440-36-0
RN
    7440-38-2
RN
    7440-42-8
RN
    7723-14-0
RN
RN
    64-17-5
RN
    67-63-0
    67-64-1
RN
    108-94-1
RN
RN
     1303-86-2
     7697-37-2
RN
     7727-37-9
RN
     30136-13-1
RN
     7631-86-9P
RN
     12125-01-8P
RN
     7440-21-3P
RN
     11105-01-4P
RN
    12033-89-5P
RN
     1344-28-1
RN
     6876-41-1D
RN
     7429-90-5
RN
     24304-00-5
RN
     153315-81-2
RN
L10 ANSWER 7 OF 16 CAPLUS COPYRIGHT 2003 ACS
     1999:412782 CAPLUS
AN
     131:65892
DN
     Manufacture of substrates with patterned polysilane layers
TΙ
     Fukushima, Motoo; Mori, Shigeru
IN
     Shin-Etsu Chemical Industry Co., Ltd., Japan
PΑ
     Jpn. Kokai Tokkyo Koho, 5 pp.
     CODEN: JKXXAF
     Patent
DT
     Japanese
LA
     ICM G03F007-075
IC
     ICS G03F007-32; G03F007-38; H05K003-00
     74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other
CC
     Reprographic Processes)
     Section cross-reference(s): 38
FAN.CNT 1
                                           APPLICATION NO. DATE
                      KIND DATE
     PATENT NO.
                            -----
                      ____
                                            JP 1997-362183 19971211
                            19990702
     JP 11174682
                       A2
PΙ
                       B2
                            20020715
     JP 3301370
                                            US 1998-208588
                                                            19981210
                      Α
                            20000829
     US 6110651
                     Α
                            19971211
PRAI JP 1997-362183
     The title process involves: (1) formation of polysilane layer on a
     substrate, (2) selective light irradn. of the layer in presence of a
     solvent which dissolves siloxane but not polysilane, (3) removal
     of siloxane by treatment in a solvent that only dissolves
     siloxane, and (4) complete removal of the solvent. Polysilane may
     be (RlmR2nXpSi)q [R1-2 = (un)substituted hydrocarbon; X = (un)substituted
     hydrocarbon, alkoxy, halogen; 1 .ltoreq. m + n + p .ltoreq. 2.2; 10
      .1toreq. q.1toreq. 100,000; q = integer]. Polysilane patterns are formed
     with high accuracy. The process is useful in manuf. of printed
      circuit boards, battery electrodes, sensors, etc.
```

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polysilane patterning siloxane dissoln
ST
    Photoimaging materials
ΙT
        (manuf. of patterned polysilane by dissoln of siloxanes by
        irradn. in presence of alc.)
     Polysilanes
ΙT
    RL: PEP (Physical, engineering or chemical process); PNU (Preparation,
     unclassified); TEM (Technical or engineered material use); PREP
     (Preparation); PROC (Process); USES (Uses)
        (manuf. of patterned polysilane by dissoln of siloxanes by
        irradn. in presence of alc.)
     Polysiloxanes, processes
ΙT
     RL: PNU (Preparation, unclassified); REM (Removal or disposal); PREP
     (Preparation); PROC (Process)
        (manuf. of patterned polysilane by dissoln of siloxanes by
        irradn. in presence of alc.)
     Alcohols, uses
IΤ
     RL: TEM (Technical or engineered material use); USES (Uses)
        (manuf. of patterned polysilane by dissoln of siloxanes by
        irradn. in presence of alc.)
     146088-00-8P, Poly(methylphenylsilane)
IT
     RL: PEP (Physical, engineering or chemical process); PNU (Preparation,
     unclassified); TEM (Technical or engineered material use); PREP
     (Preparation); PROC (Process); USES (Uses)
        (manuf. of patterned polysilane by dissoln of siloxanes by
        irradn. in presence of alc.)
     64-17-5, Ethanol, uses 67-63-0, Isopropanol, uses
ΙT
     RL: TEM (Technical or engineered material use); USES (Uses)
        (manuf. of patterned polysilane by dissoln of siloxanes by
        irradn. in presence of alc.)
     146088-00-8P
RN
     64-17-5
RN
     67-63-0
RN
L10 ANSWER 8 OF 16 CAPLUS COPYRIGHT 2003 ACS
     1976:6867 CAPLUS
AN
     84:6867
DN
     Detergent for cleaning a polymer surface
ΤI
     Bratolyubov, A. S.; Rýbak, I. K.; Grabovskii, O. Z.; Bykov, M. V.;
ΙN
     Goncharenko, G. S.; Lugina, V. P.; Finogeev, N. D.
PΑ
     USSR
     U.S.S.R.
SO
     From: Otkrytiya, Izobret., Prom. Obraztsy, Tovarnye Znaki 1975, 52(36),
     CODEN: URXXAF
 DΤ
     Patent
     Russian
 A_{i,I}
 T.C.
     C11D
      46-6 (Surface Active Agents and Detergents)
 CC
 FAN.CNT 1
                                           APPLICATION NO. DATE
      PATENT NO.
                     KIND DATE
                                           _____
      ______
                                           SU 1974-2003326 19740311
                      T 19750930
      SU 486040
 PRAI SU 1974-2003326
                           19740311
     Mixts. of Cl2FCCClF2 [76-13-1] 35-45, MeCCl3 [71-55-6] 10-20, and aliph.
      alc. (EtOH [64-17-5] and/or iso-PrOH [67-63-0]) 40-50 vol. were
      useful for cleaning polymer surfaces (e.g., elec. circuit
      boords) with min. damage to siloxane elec. insulation.
      detergent cleaning circuit board; elec circuit board
 ST
      cleaning; solvent cleaning circuit board; haloalkane cleaning
      circuit board; alc cleaning circuit board
      Detergents
 ΙT
         (cleaning solvents, for soldering flux removal from printed
```

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circuits)
    Electric circuits
IΤ
        (printed, soldering flux removal from, solvents for)
    64-17-5, uses and miscellaneous 67-63-0, uses and miscellaneous
IT
             76-13-1
     71-55-6
    RL: TEM (Technical or engineered material use); USES (Uses)
        (cleaning compns. contg., for elec. circuit boards)
     64-17-5
RN
     67-63-0
RN
     71-55-€
RN
RN
    76-13-1
L10 ANSWER 9 OF 16 CAPLUS COPYRIGHT 2003 ACS
     2000:381447 CAPLUS
AN
     132:355738
DN
    Method of reducing spiral defects in glass by adding an isopropyl alcohol
ΤT
     rinse step before depositing spin-on glass in integrated-circuit
     fabrication
     Chiang, Chen-Chia; Lin, Chung-An
IN
     Taiwan Semiconductor Manufacturing Company, Taiwan
PA
     U.S., 8 pp.
SO
     CODEN: USXXAM
     Patent
DT
     English
LA
     ICM H01L021-31
TC
     ICS H01L021-469
NCL 438778000
     76-3 (Electric Phenomena)
CC
     Section cross-reference(s): 57, 66
FAN.CNT 1
                                           APPLICATION NO. DATE
                     KIND DATE
     PATENT NO.
                                            _____
     _____
                      ____
                                            US 1998-135042
                                                            19980817
   US 6071831
                            20000606
                       Α
PΙ
                            19980817
PRAI US 1998-135042
     A method of forming an interlevel dielec. layer of spin-on-glass is
     described which avoids spiral defects from occurring in the layer of
     spin-on-glass. Before the spin-on-glass is deposited and with the wafer
     spinning at a low angular velocity a 1st vol. of iso-Pr alc. is deposited
     on the wafer. The wafer continues to spin at the low angular velocity for a short time. With the wafer continuing to spin at the low angular
     velocity a 2nd vol., less than the 1st vol., of spin-on-glass is deposited
     on the wafer. The wafer continues to spin at the low angular velocity for
     a short time and then is spun at a high angular velocity for a longer
     time. The wafer is then removed from the app. used to deposit the
     spin-on-glass and processing of the wafer continues. Spiral defects in
     the layer of spin-on-glass are avoided.
     spin on glass spiral defect isopropanol rinsing silicon; integrated
     circuit rotation spin glass coating
     Polysiloxanes, processes
ΤТ
     RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical
     process); PROC (Process); USES (Uses)
         (Me; method of reducing spiral defects in glass by adding iso-Pr alc.
         rinse step for silicon before depositing spin-on glass in integrated-
         circuit fabrication)
      Dielectric films
      Integrated circuits
      Rotation
      Semiconductor device fabrication
         (method of reducing spiral defects in glass by adding iso-Pr alc. rinse
         step for silicon before depositing spin-on glass in integrated-
         circuit fabrication)
     Glass, properties
 ΙT
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RL: DEV (Device component use); PEP (Physical, engineering or chemical
    process); PRP (Properties); PROC (Process); USES (Uses)
       (spin on; method of reducing spiral defects in glass by adding iso-Pr
       alc. rinse step for silicon before depositing spin-on glass in
       integrated-circuit fabrication)
ΙT
    Coating process
    Washing
       (spin; method of reducing spiral defects in glass by adding iso-Pr alc.
       rinse step for silicon before depositing spin-on glass in integrated-
       circuit fabrication)
    7440-21-3, Silicon, processes
IT
    RL: DEV (Device component use); PEP (Physical, engineering or chemical
     process); PROC (Process); USES (Uses)
       (method of reducing spiral defects in glass by adding iso-Pr alc. rinse
       step for silicon before depositing spin-on glass in integrated-
       circuit fabrication)
     64-17-5, Ethanol, processes 67-63-0, Isopropyl alcohol,
TΨ
     processes 71-36-3, 1-Butanol, processes
     RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical
     process); PROC (Process); USES (Uses)
        (method of reducing spiral defects in glass by adding iso-Pr alc. rinse
       step for silicon before depositing spin-on glass in integrated-
       circuit fabrication)
             THERE ARE 14 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT 14
RE
(1) Akram; US 5925410 1999
(2) Chang; ULSI Technology 1996, P92 CAPLUS
(3) Gardner; US 5535525 1996
(4) Ikeno; US 5264246 1993
(5) Ilg; US 5807792 1998 CAPLUS
(6) Kim; US 5874128 1999
(7) Lin; US 5646071 1997 CAPLUS
(8) Sanada; US 5843527 1998
(9) Sato; US 6022806 2000 CAPLUS
(10) Shirley; US 5912049 1999
(11) Wang; US 5780105 1998
(12) Wilson; US 5990014 1999 CAPLUS
(13) Yen; US 6004622 1999 CAPLUS
(14) Ziger; US 5272118 1993
RN 7440-21-3
RN 64-17-5
RN 67-63-0
RN 71-36-3
L10 ANSWER 10 OF 16 CAPLUS COPYRIGHT 2003 ACS
    2000:861954 CAPLUS
AN
    134:20313
DN
    Vapor deposition for protective coating on spherical solder-alloy powders
ΤŢ
IN Minoque, Gerard R.
    Alpha Metals, Inc., USA
PΑ
     PCT Int. Appl., 16 pp.
SO
     CODEN: PIXXD2
     Patent
DΤ
     English
LΑ
     ICM H01L021-48
TC
     ICS H01L023-498; H05K003-34; B23K035-02
     56-9 (Nonferrous Metals and Alloys)
     Section cross-reference(s): 76
 FAN.CNT 1
                                      APPLICATION NO. DATE
     PATENT NO. KIND DATE
                                          _____
      ______
    WO 2000074132 A1 20001207
                                         WO 2000-US15220 20000601
 PT
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AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR,
            CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU,
            ID, IL, IN, IS, JP, KE, KG, KP, KR, K2, LC, LK, LR, LS, LT, LU,
            LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD,
            SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU,
            ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
        RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
            DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ,
            CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
                                                            20000531
                                          US 2000-584674
                            20030114
                      Bl
    US 6506448
                                                            20000601
                                           EP 2000-938067
                            20020821
                      Α1
     EP 1232524
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
            IE, SI, LT, LV, FI, RO, MK, CY, AL
                            19990601
PRAI US 1999-137031P P
                      A2
                            20000531
     us 2000-584674
                            20000601
     WO 2000-US15220
                      W
     The surface of spherical solder-alloy powder is coated by vapor deposition
AB
    using a soln. contg.: (a) volatile org. solvent; (b) low-viscosity oil or
    org. acid at nominally 0.05-5.0+; and (c) a surfactant at 0.01-1.0+, and
     optional flux and/or UV-fluorescent dye. The solder-alloy powder in a
     chamber is dipped into the coating soln. in a vapor-tight 2nd chamber, and
     after a predetd. time is removed, placed into 3rd vapor-tight low-pressure
     chamber, and heated to 54-121.degree. to vaporize the residual solvent
    with optional recovery. The coated alloy powder can be stored without
     corrosion, and can be used in elec. conducting solder pastes or adhesives
     The coated solder spheres are suitable for assembly of elec.
     circuit boards with surface-mounted electronic parts, followed by
     heating for reflow soldering. The typical coating soln. contains AcOH as
     the solvent, paraffin oil or isostearic acid as the low-viscosity addn.,
     simethicone as the surfactant, and a polar or nonpolar flux.
     solder alloy ball protective coating soln; elec circuit assembly
ST
     coated solder ball
     Fluorescent dyes
ΤT
        (UV-, coating contg.; soln. for vapor deposition of protective coating
        on spherical solder-alloy powders)
ΙT
     Electric circuits
        (assemblies, spherical solder for; soln. for vapor deposition of
        protective coating on spherical solder-alloy powders)
     Lubricating oils
IT
     Surfactants
        (coating contg.; soln. for vapor deposition of protective coating on
        spherical solder-alloy powders)
     Paraffin oils
IΤ
       Polysiloxanes, uses
     RL: MOA (Modifier or additive use); USES (Uses)
        (coating contg.; soln. for vapor deposition of protective coating on
        spherical solder-alloy powders)
     Cyclosiloxanes
ΙT
     RL: MOA (Modifier or additive use); USES (Uses)
        (di-Me, surfactant, coating bath contg.; soln. for vapor deposition of
        protective coating on spherical solder-alloy powders)
     Polyolefins
ΙT
     RL: MOA (Modifier or additive use); USES (Uses)
        (oil, coating contg.; soln. for vapor deposition of protective coating
        on spherical solder-alloy powders)
     Solders
IT
        (spheres, coating of; soln. for vapor deposition of protective coating
        on spherical solder-alloy powders)
     64-17-5, Ethanol, uses 67-63-0, Isopropanol, uses
                                                          67-64-1,
ΙT
     Acetone, uses 79-01-6, Trichloroethylene, uses 106-94-5, n-Propyl
               124-04-9, Adipic acid, uses
                                             30399-84-9, Isostearic acid
     RL: MOA (Modifier or additive use); USES (Uses)
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(coating bath contg.; soln. for vapor deposition of protective coating on spherical solder-alloy powders) 8050-81-5, Simethicone 541-02-6, Decamethylcyclopentasiloxane ΙT RL: MOA (Modifier or additive use); USES (Uses) (surfactant, coating bath contg.; soln. for vapor deposition of protective coating on spherical solder-alloy powders) THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD RE.CNT RF. (1) Fry Metals Inc; WO 9701414 A 1997 CAPLUS (2) Fujitsu Ltd; EP 0539211 A 1993 (3) Murata Manufacturing Co; EP 0905775 A 1999 (4) Union Carbide Chem Plastic; EP 0556864 A 1993 CAPLUS 64 - 17 - 5RN 67-63-0 RN 67-64-1 RN79-01-6 RN 106-94-5 RN 124-04-9 RN 30399-84-9 RN 541-02-6 RN 8050-81-5 RN L10 ANSWER 11 OF 16 CAPLUS COPYRIGHT 2003 ACS 1996:437814 CAPLUS ΑN 125:89676 DN Cleaning agent, method and equipment TIKumagai, Masaru; Shimozawa, Hiroshi; Oguni, Naoyuki; Inada, Minoru; Saito, ΤN Nobuhiro; Yamafuji, Shigeo; Shimizu, Chiyuki; Umehara, Kazunori Kabushiki Kaisha Toshiba, Japan; Toshiba Silicone Co., Ltd. PΑ PCT Int. Appl., 124 pp. SO CODEN: PIXXD2 Patent DTJapanese LA ICM B08B003-08 IC 46-6 (Surface Active Agents and Detergents) Section cross-reference(s): 76 FAN.CNT 1 APPLICATION NO. DATE KIND DATE PATENT NO. _____ _ _ _ _ _____ WO 1995-JP2149 19951019 A1 19960502 WO 9612571 PΙ W: CN, JP, KR, US RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE EP 1995-934846 19951019 A1 19970806 EP 787537 R: DE, ES, FR, GB, IT, NL A 19980114 19951019 CN 1995-196867 CN 1170374 JP 1996-513771 19951019 B2 20000124 JP 3002261 A 19941019 PRAI JP 1994-253487 JP 1994-253488 Α 19941019 19941019 JP 1994-253489 A JP 1994-253490 A 19941019 A 19941019 JP 1994-253491 A 19941019 JP 1994-253492 A 19950410 JP 1995-84066 JP 1995-108410 A 19950502 JP 1995-108411 A 19950502 A 19950502 W 19951019 JP 1995-108412 WO 1995-JP2149 The title cleaning agent can substitute for chlorofluorocarbon or AΒ chlorinated solvents. In particular, the method is for cleaning electronics parts such as printed boards or mounted parts or metallic parts, and is possible to remove various contaminants from the parts and dry the same to the extent comparative to that of the method of cleaning

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with chlorofluorocarbon or chlorinated solvents. Examples of the cleaning
agent include polar cleaning agents with a soly. parameter of .gtoreq.9 or
a permittivity of .gtoreq.4, mixts. thereof with other cleaning agents
with a soly. parameter of <9 and a permittivity of <4, and cleaning agents
contg. as the active ingredient an azeotropic or pseudoazeotropic compn.
contg. a low-mol.-wt. siloxane compd. The cleaning method is
characterized by comprising .gtoreq.1 step selected from the group
consisting of the step of cleaning the object with a polar cleaning agent
having a soly. parameter of .gtoreq.9 or a permittivity of .gtoreq.4, the
step of rinsing the object with a mixt. of the above polar cleaning agent
with another cleaning agent having a soly. parameter of <9 and a
permittivity of <4, and the drying step.
polar cleaning agent electronic part; azeotropic siloxane
cleaning compn; pseudoazeotropic siloxane cleaning compn;
chlorofluorocarbon substitute cleaning agent; chlorinated solvent
substitute cleaning agent; fluorocarbon cleaning agent electronic part;
water drying agent app
Drying apparatus
Electric apparatus
   (cleaning agent, method and equipment for electron parts)
Hydrocarbons, uses
RL: PRP (Properties); TEM (Technical or engineered material use); USES
   (cleaning agent, method and equipment for electron parts)
Acetals
Alcohols, uses
Amides, uses
Amines, uses
Anhydrides
  Cyclosiloxanes
Esters, uses
Ethers, uses
Glycols, uses
Ketones, uses
Nitriles, uses
Perfluorocarbons
Phenols, uses
Quaternary ammonium compounds, uses
Sulfoxides
RL: TEM (Technical or engineered material use); USES (Uses)
    (cleaning agent, method and equipment for electron parts)
Siloxanes and Silicones, uses
RL: TEM (Technical or engineered material use); USES (Uses)
    (low-mol.-wt.; cleaning agent, method and equipment for electron parts)
Degreasing
    (agents, cleaning agent, method and equipment for electron parts)
Cleaning
    (app., cleaning agent, method and equipment for electron parts)
Detergents
   (azeotropic, cleaning agent, method and equipment for electron parts)
Hydrocarbons, uses
RL: TEM (Technical or engineered material use); USES (Uses)
    (chloro fluoro, cleaning agent, method and equipment for electron
   parts)
 Detergents
    (cleaning compns., cleaning agent, method and equipment for electron
    parts)
 Detergents
    (cleaning compns., spray, cleaning agent, method and equipment for
    electron parts)
 Hydrocarbons, uses
 RL: PRP (Properties); TEM (Technical or engineered material use); USES
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ST

ΙT

TΤ

TΤ

IT

IΤ

ΙΤ

ΙT

IT

ΙT

ΙT

IΤ

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(fluoro, cleaning agent, method and equipment for electron parts)
     Alcohols, uses
ΙT
     RL: TEM (Technical or engineered material use); USES (Uses)
        (fluoro, cleaning agent, method and equipment for electron parts)
     Perhalocarbons
ΙT
     RL: TEM (Technical or engineered material use); USES (Uses)
        (fluoroiodo, cleaning agent, method and equipment for electron parts)
     Electric circuits
ΙT
        (printed, boards, cleaning agent, method and equipment for electron
        parts)
                                                             178820-78-5
                                              178820-77-4
                   178820-75-2 178820-76-3
     178820-74-1
ΙT
     178820-79-6 178820-80-9
                               178820-81-0
     RL: PRP (Properties); TEM (Technical or engineered material use); USES
        (azeotropic; cleaning agent, method and equipment for electron parts)
     64-17-5, Ethanol, uses 64-19-7, Acetic acid, uses 67-63-0,
IΤ
     2-Propanol, uses 67-64-1, Acetone, uses 71-23-8, 1-Propanol, uses
     75-05-8, Acetonitrile, uses 75-52-5, Nitromethane, uses 75-65-0, tert-Butanol, uses 75-85-4, tert-Pentyl alcohol 75-89-8 76-05-
                                                                    76-05-1,
     Trifluoroacetic acid, uses 78-82-0, Isobutyronitrile 78-93-3, Methyl
     ethyl ketone, uses 79-24-3, Nitroethane 96-22-0, 3-Pentanone
     105-37-3, Ethyl propionate 107-12-0, Propionitrile
                                                            107-15-3,
     Ethylenediamine, uses 107-46-0, Hexamethyldisiloxane
     107-51-7, Octamethyltrisiloxane 108-21-4, Isopropyl acetate
     109-60-4, Propyl acetate 109-73-9, Butylamine, uses 109-74-0,
     Butyronitrile 109-86-4, 2-Methoxyethanol 109-99-9, THF, uses
                          110-71-4, 1,2-Dimethoxyethane 110-85-0,
     110-01-0, Thiophane
     Piperazine, uses 110-91-8, Morpholine, uses 111-27-3, 1-Hexanol, uses
     112-34-5, Diethylene glycol monobutyl ether 123-51-3, Isopentyl alcohol
     123-86-4, Butyl acetate 138-22-7, Butyl lactate
                                                        141-78-6, Ethyl
                    141-79-7, Mesityl oxide 307-34-6
                                                         335-57-9, Fluorinert
     acetate, uses
               352-93-2, Diethyl sulfide 355-42-0 378-94-9,
     PF 5070
                                      422-56-0 542-55-2, Isobutyl formate
     Perfluoromorpholine 382-28-5
                                               583-59-5,
     556-67-2, Octamethylcyclotetrasiloxane
                                                  872-50-4, NMP, uses
     2-Methylcyclohexanol 591-78-6, 2-Hexanone
                 3452-97-9, 3,5,5-Trimethylhexanol 7664-38-2, Phosphoric
     1320-67-8
                  51000-94-3 52125-53-8, Propylene glycol monoethyl ether
     acid, uses
     56539-66-3, 3-Methoxy-3-methyl-1-butanol 75330-23-3 111109-77-4,
     Dipropylene glycol dimethyl ether 139063-93-7
     RL: PRP (Properties); TEM (Technical or engineered material use); USES
         (cleaning agent, method and equipment for electron parts)
     178820-74-1
 RN
     178820-75-2
 RN
 RN
     178820-76-3
     178820-77-4
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     178820-78-5
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     178820-79-6
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     178820-80-9
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     178820-81-0
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      64-17-5
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      64-19-7
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      67-64-1
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      75-05-8
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      75-52-5
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      75-65-0
 RN
      75-85-4
 RN
      75-89-8
 RN
      76-05-1
 RN
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(Uses)

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78-82-0
RN
     78-93-3
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     79-24-3
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     96-22-0
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     105-37-3
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RN
     107-12-0
     107-15-3
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     107-46-0
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     107-51-7
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     108-21-4
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     109-60-4
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     109-73-9
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     109-74-0
     109-86-4
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     109-99-9
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     110-01-0
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     352-93-2
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     355-42-0
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     378-94-9
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     382-28-5
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     422-56-0
     542-55-2
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     556-67-2
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      583-59-5
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      591-78-6
RN
      872-50-4
      1320-67-8
RN
      3452-97-9
RN
      7664-38-2
RN
      51000-94-3
RN
      52125-53-8
RN
      56539-66-3
RN
      75330-23-3
RN
      111109-77-4
RN
      139063-93-7
RN
    ANSWER 12 OF 16 CAPLUS COPYRIGHT 2003 ACS
L10
      2001:731251 CAPLUS
AN
      135:281843
 DN
      Method of producing a multilayered wiring board with small area via holes
 TΙ
      Toyoshima, Toshiyuki; Yanaura, Satoshi; Furuhashi, Yasuo; Fujioka,
 ΙN
      Hirofumi
 PA
      Japan
      U.S. Pat. Appl. Publ., 18 pp.
 SO
      CODEN: USXXCO
 DT
      Patent
      English
 LΑ
      ICM H05K003-02
 IC
      ICS H05K003-10
     029852000
 NCL
```

76-14 (Electric Phenomena) CC Section cross-reference(s): 38 FAN.CNT 1 APPLICATION NO. DATE PATENT NO. KIND DATE _____ _____ ____ US 2000-738855 20001218 US 2001025414 A1 20011004 PΙ JP 2000-97250 20000331 20011012 JP 2001284813 A2 20000331 PRAI JP 2000-97250 А Method of producing a multilayered wiring board comprising the steps of subjecting the photosensitive resin to exposure- and development-treatment to form the holes having a predetd. size and shape; depositing and forming the curable resin to the insulating layer having the holes formed therein in such a manner as to bury the holes, and conducting heat-treatment to form the cured thin film of the curable resin on the surface of the insulating layer; and so removing the curable resin as to leave the cured thin film to obtain the via-holes having the reduced opening size by the cured thin film. printed circuit board via hole polymer resin STPolyvinyl acetals ΙT RL: PEP (Physical, engineering or chemical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses) (S-Lec KW 3; method of producing multilayered wiring board with small \langle area via holes) Epoxy resins, processes IT RL: PEP (Physical, engineering or chemical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses) (acrylates; method of producing multilayered wiring board with small area via holes) Coating process ΙT Contact holes Crosslinking Dielectric films Heat treatment Printed circuit boards (method of producing multilayered wiring board with small area via holes) ITAcrylic polymers, processes Aminoplasts Butadiene rubber, processes Phenolic resins, processes Polysiloxanes, processes RL: PEP (Physical, engineering or chemical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses) (method of producing multilayered wiring board with small area via holes) Epoxy resins, processes IΤ RL: PEP (Physical, engineering or chemical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses) (photosensitive; method of producing multilayered wiring board with small area via holes) Interconnections (electric) TΤ (vias; method of producing multilayered wiring board with small area via holes) 25068-38-6 IT RL: PEP (Physical, engineering or chemical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses) (E 1001; method of producing multilayered wiring board with small area via holes) 9003-17-2 TΤ

RL: PEP (Physical, engineering or chemical process); TEM (Technical or

(butadiene rubber, method of producing multilayered wiring board with

engineered material use); PROC (Process); USES (Uses)

```
small area via holes)
    362660-00-2, CPC 8000
ΙT
    RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical
    process); PROC (Process); USES (Uses)
        (copper paste; method of producing multilayered wiring board with small
        area via holes)
     1310-73-2, Sodium hydroxide, processes
ΙΤ
     RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical
     process); PROC (Process); USES (Uses)
        (developer; method of producing multilayered wiring board with small
        area via holes)
     67-63-0, Isopropanol, processes 111-76-2, Butyl cellosolve
ΙT
     7738-94-5, Chromic acid (H2CrO4)
     RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical
     process); PROC (Process); USES (Uses)
        (method of producing multilayered wiring board with small area via
        holes)
                                              621-82-9D, Cinnamic acid, esters
     471-34-1, Calcium carbonate, processes
IT
                       9003-08-1, Cymel 370
                                                9003-18-3, Butadiene-
     2669-72-9, MX 280
                              58607-87-7 71868-10-5, Irgacure 907
     acrylonitrile copolymer
     115166-29-5, Epikote 180 210106-42-6, XP 9500CC
                                                         362660-64-8, JSR-KS 22
     362660-65-9, Probelec XB 7081
     RL: PEP (Physical, engineering or chemical process); TEM (Technical or
     engineered material use); PROC (Process); USES (Uses)
        (method of producing multilayered wiring board with small area via
     362658-73-9, OM 950
IT
     RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical
     process); PROC (Process); USES (Uses)
        (neutralizing agent; method of producing multilayered wiring board with
        small area via holes)
     25068-38-6
RN
     9003-17-2
RN
     362660-00-2
RN
RN
     1310-73-2
RN
     67-63-0
RN
     111-76-2
     7738-94-5
RN
     471-34-1
RN
     621-82-9D
RN
     2669-72-9
RN
     9003-08-1
RN
     9003-18-3
RN
     58607-87-7
RN
RN
     71868-10-5
     115166-29-5
RN
     210106-42-6
RN
     362660-64-8
RN
     362660-65-9
RN
     362658-73-9
RN
L10 ANSWER 13 OF 16 CAPLUS COPYRIGHT 2003 ACS
     1999:650669 CAPLUS
AN
     131:275831
DN
     Automated detection and reporting of volatile organic compounds (VOCs) in
ΤI
     complex environments
     Hargis, P. J., Jr.; Preppernau, B. L.; Osbourn, G. C.; Ricco, A. J.; Frye,
ΑU
     G. C.
     Laser, Optics and Remote Sensing Department, Sandia National Laboratories,
 CS
     Albuquerque, NM, 87185-1423, USA
     Sandia National Laboratories [Technical Report] SAND (1997), SAND97-0509,
 SO
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i-vii, 1-18

CODEN: SNLSDT

DT Report

LA English

CC 59-1 (Air Pollution and Industrial Hygiene)

Section cross-reference(s): 47, 80

Results of work to develop volatile org. compd. (VOC) sensing systems AΒ based on 2 complementary techniques are discussed. The first technique used a gated channeltron detector for resonant laser-induced multiphoton photoionization detection of trace org. vapors in a supersonic mol. beam. The channeltron was gated using a relatively simple circuit to generate a neg. gate pulse with a width of 400 ns (FWHM), a 50 ns turn-on (rise) time, a 1.5 .mu.s turn-off (decay) time, a pulse amplitude of -1000 V, and a DC offset adjustable from 0 to -1500 V. This gated channeltron rejects spurious responses to UV laser light scattered directly into the channeltron and time-delayed ionization signals induced by photoionization of residual gas in the vacuum chamber. Detection limits in the part-per-trillion range were demonstrated with the gated detector. second technique used arrays of surface acoustic wave (SAW) devices coated with various chem. selective materials (e.g., polymers, self-assembled monolayers) to provide unique response patterns to various chem. analytes. This work focused on polymers, formed by spin casting from soln. or by plasma polymn., as well as on self-assembled monolayers. Response from coated SAW to various concns. of water, VOC, and organophosphonates (chem. warfare agent simulants) were used for calibration data. A novel visual empirical region of influence (VERI) pattern recognition technique was used to evaluate the ability to use these response patterns to correctly identify chem. species. This study showed how the VERI technique can be used to det. the best set of coatings materials, and to identify unknown analytes based on previous calibration data.

volatile org detn monitoring air analysis; chem sensor automated volatile org detn monitoring; laser induced multiphoton ionization volatile org

detection; surface acoustic wave sensor volatile org detection

IT Air analysis

Sensors

(automated volatile org. compd. and mixt. detn. and monitoring in air using laser-induced multiphoton ionization and surface acoustic wave microsensor array-equipped chem. sensors)

IT Volatile organic compounds

RL: ANT (Analyte); POL (Pollutant); ANST (Analytical study); OCCU (Occurrence)

(automated volatile org. compd. and mixt. detn. and monitoring in air using laser-induced multiphoton ionization and surface acoustic wave microsensor array-equipped chem. sensors)

IT Polysiloxanes, uses

RL: DEV (Device component use); TEM (Technical or engineered material use); USES (Uses)

(cyanide-modified; sensor coating; automated volatile org. compd. and mixt. detn. and monitoring in air using laser-induced multiphoton ionization and surface acoustic wave microsensor array-equipped chem. sensors)

IT Polyoxyalkylenes, uses

RL: DEV (Device component use); TEM (Technical or engineered material use); USES (Uses)

(fluorine- and sulfo-contg., ionomers, sensor coating; automated volatile org. compd. and mixt. detn. and monitoring in air using laser-induced multiphoton ionization and surface acoustic wave microsensor array-equipped chem. sensors)

IT Polyoxyalkylenes, uses

RL: DEV (Device component use); TEM (Technical or engineered material use); USES (Uses)

(fluorine-contg., sulfo-contg., ionomers, sensor coating; automated volatile org. compd. and mixt. detn. and monitoring in air using

laser-induced multiphoton ionization and surface acoustic wave microsensor array-equipped chem. sensors)

IT Alcohols, uses

RL: DEV (Device component use); TEM (Technical or engineered material use); USES (Uses)

(polyhydric, fluoro, sensor coating; automated volatile org. compd. and mixt. detn. and monitoring in air using laser-induced multiphoton ionization and surface acoustic wave microsensor array-equipped chem. sensors)

IT Fluoropolymers, uses Fluoropolymers, uses

RL: DEV (Device component use); TEM (Technical or engineered material use); USES (Uses)

(polyoxyalkylene-, sulfo-contg., ionomers, sensor coating; automated volatile org. compd. and mixt. detn. and monitoring in air using laser-induced multiphoton ionization and surface acoustic wave microsensor array-equipped chem. sensors)

IT Ionomers

RL: DEV (Device component use); TEM (Technical or engineered material use); USES (Uses)

(polyoxyalkylenes, fluorine- and sulfo-contg., sensor coating; automated volatile org. compd. and mixt. detn. and monitoring in air using laser-induced multiphoton ionization and surface acoustic wave microsensor array-equipped chem. sensors)

IT 56-23-5, analysis 67-56-1, Methanol, analysis 67-63-0, Isopropanol, analysis 67-64-1, 2-Propanone, analysis 67-66-3, Chloroform, analysis 71-23-8, n-Propanol, analysis 71-43-2, Benzene, analysis 78-93-3, Methyl ethyl ketone, analysis 79-01-6, Trichloroethylene, analysis 108-88-3, Toluene, analysis 110-54-3, Hexane, analysis 110-82-7, Cyclohexane, analysis 127-18-4, Perchloroethylene, analysis 464-07-3, Pinacolyl alcohol 756-79-6, Dimethyl methyl phosphonate 1445-75-6, Diisopropyl methyl phosphonate 5989-27-5 26635-64-3, Isooctane

RL: ANT (Analyte); POL (Pollutant); ANST (Analytical study); OCCU (Occurrence)

(automated volatile org. compd. and mixt. detn. and monitoring in air using laser-induced multiphoton ionization and surface acoustic wave microsensor array-equipped chem. sensors)

IT 24937-05-1, Polyethylene glycol adipate

RL: DEV (Device component use); TEM (Technical or engineered material use); USES (Uses)

(cyanide-modified; sensor coating; automated volatile org. compd. and mixt. detn. and monitoring in air using laser-induced multiphoton ionization and surface acoustic wave microsensor array-equipped chem. sensors)

TT 79-10-7, 2-Propenoic acid, uses
RL: DEV (Device component use); TEM (Technical or engineered material use); USES (Uses)

(plasma-grafted; sensor coating; automated volatile org. compd. and mixt. detn. and monitoring in air using laser-induced multiphoton ionization and surface acoustic wave microsensor array-equipped chem. sensors)

97-53-0, Eugenol 9002-98-6, Poly(ethylenimine) 9003-20-7, Poly(vinyl acetate) 9003-27-4, Polyisobutylene 9004-57-3, Ethyl cellulose 24969-06-0, Poly(epichlorohydrin) 25035-84-1, Poly(vinyl propionate) 25791-89-3 28212-48-8, Poly(diphenoxy phosphazene) 135311-44-3 RL: DEV (Device component use); TEM (Technical or engineered material use); USES (Uses)

(sensor coating; automated volatile org. compd. and mixt. detn. and monitoring in air using laser-induced multiphoton ionization and surface acoustic wave microsensor array-equipped chem. sensors)

RE.CNT 14 THERE ARE 14 CITED REFERENCES AVAILABLE FOR THIS RECORD

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    1994, P180
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     56-23-5
RN
     67-56-1
     67-63-0
RN
     67-64-1
RN
     67-66-3
RN
     71-23-8
RN
     71-43-2
RN
     78-93-3
RN
     79-01-6
RN
     108-88-3
RN
     110-54-3
RN
RN
     110-82-7
     127-18-4
RN
RN
     464-07-3
RN
     756-79-6
     1445-75-6
RN
     5989-27-5
RN
     26635-64-3
RN
     24937-05-1
RN
RN
     79-10-7
     97-53-0
RN
     9002-98-6
RN
     9003-20-7
RN
     9003-27-4
RN
     9004-57-3
RN
RN
     24969-06-0
RN
     25035-84-1
     25791-89-3
RN
     28212-48-8
RN
     135311-44-3
RN
L10 ANSWER 14 OF 16 CAPLUS COPYRIGHT 2003 ACS
     1998:282379 CAPLUS
AN
DN
     128:329869
     Technique for the removal of residual spin-on-glass (SOG) after full SOG
TΙ
     Wu, Lin-june; Yu, Chen-hua Douglas; Lee, Jin-yuan
ΙN
     Taiwan Semiconductor Manufacturing Company, Ltd., Taiwan
PA
     U.S., 9 pp.
SO
     CODEN: USXXAM
DT
     Patent
     English
LA
     ICM H01L021-4763
ΙC
NCL
     438624000
     76-3 (Electric Phenomena)
CC
FAN.CNT 1
```

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 5747381	A	19980505 19960212	US 1996-599770	19960212
PRAI AB	This invention relates to a method for removing residual spin-on-glass (SOG) during a planarization processing step wherein the SOG is used as a sacrificial planarization medium and subjected to a full etchback to an underlying interlevel dielec. (ILD) layer. The SOG is applied over the ILD layer, and etched back into the ILD layer by reactive-ion-etching under conditions of comparable etch rates for both SOG and ILD. At endpoint there some residual pockets of SOG can be present as well as a region of SOG along the edges of the wafer where it is clamped in the etchback tool. The residual SOG must be removed completely to avoid SOG cracking after thermal processing and SOG outgassing during subsequent metal deposition. For this purpose an aq. etch consisting of hydrofluoric acid buffered with ammonium fluoride is used. The etchant compn. chosen exhibits a selectivity for SOG over the ILD glass of greater than 40 making it suitable for removing considerable SOG residues with minimal attack of the ILD.				
ST	spin on glass removal planarization IC				
IT	Etching (anisotropic; technique for removal of residual spin-on-glass (SOG) after full SOG etchback)				
IΤ					
11	(etching, reactive; technique for removal of residual spin-on-glass (SOG) after full SOG etchback)				
IT	Glass, processes RL: PEP (Physica	il, eng	ineering or c	hemical process); [REM (Removal or
	full SOG etch	chnique	for removal	of residual spin-on	n-glass (SOG) after
ΙΤ	Etching (sputter, reactive; technique for removal of residual spin-on-glass (SOG) after full SOG etchback)				
ΙT	(technique fo etchback)	al, eng or remo	ineering or c	hemical process); lal spin-on-glass (PROC (Process) SOG) after full SOG
ΙΤ	dienosalle PROC	npolyme al, eng (Proce or remo	ineering or c	hemical process); lal spin-on-glass (REM (Removal or SOG) after full SOG
ΙΤ	Integrated circ MOSFET (transis Semiconductor do (technique for etchback for	uits cors) evice f or remo	abrication oval of residu	al spin-on-glass (SOG) after full SOG
ΙT		s or remo	oval of residu	al spin-on-glass (SOG) formed on an ILD
	layer)				
IT	Phosphosilicate RL: PEP (Physicate) (technique for layer contg.	al, eng or remo	rineering or o	chemical process); al spin-on-glass (PROC (Process) SOG) formed on an ILD
ΙT	67-63-0, Isopro RL: NUU (Other (technique f	pyl ald	classified);	USES (Uses) nal spin-on-glass (SOG) after full SOG
ΙT	etchback) 7631-86-9, Sili RL: PEP (Physic	ca, pro al, eno	ocesses gineering or o	chemical process);	PROC (Process)

```
technique for removal of residual spin-on-glass (SOG) formed on an ILD
       layer contg.)
     12125-01-8, Ammonium fluoride
IT
     RL: NUU (Other use, unclassified); USES (Uses)
        (technique for removal of residual spin-on-glass (SOG) hydrofluoric
       acid buffered with)
     75-73-0, Tetrafluoromethane
IT
     RL: NUU (Other use, unclassified); USES (Uses)
        (technique for removal of residual spin-on-glass (SOG) with)
     7664-39-3, Hydrofluoric acid, uses
ΙT
     RL: NUU (Other use, unclassified); USES (Uses)
        (technique for removal of residual spin-on-glass (SOG) with ammonium
        fluoride-buffered)
     18130-74-0, Bifluoride (HF21-)
ΙT
     RL: MSC (Miscellaneous)
        (technique for removal of residual spin-on-glass (SOG) with ammonium
        fluoride-buffered hydrofluoric acid)
              THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT
RE
(1) Leong; US 5192697 1993 CAPLUS
(2) Matsuura; US 5459105 1995 CAPLUS
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     67-63-0
RN
     7631-86-9
RN
     12125-01-8
RN
RN
     75-73-0
RN
     7664-39-3
     18130-74-0
RN
L10 ANSWER 15 OF 16 CAPLUS COPYRIGHT 2003 ACS
     1999:100665 CAPLUS
AN
     130:126985
DN
     Fluid property investigation by impedance characterization of quartz
TΤ
     crystal resonators. Part I: Methodology, crystal screening, and Newtonian
     fluids
     Nwankwo, E.; Durning, C. J.
ΑU
     DuPont Experimental Station, Advanced process Control and Optimization
CS
     Group, Wilmington, DE, 19880-0101, USA
     Sensors and Actuators, A: Physical (1999), A72(2), 99-109
SO
     CODEN: SAAPEB; ISSN: 0924-4247
     Elsevier Science S.A.
PΒ
     Journal
DΤ
     English
LA
     48-11 (Unit Operations and Processes)
CC
     Section cross-reference(s): 37, 76
     In the first of this two-part communication, we present a methodol. for
AΒ
     the application of thickness shear mode (TSM) quartz crystal resonators
     (QCR) in fluid property investigation. To this end, we outline a protocol
     for the prepn. of crystal surfaces for fluid contact and establish a
     methodol. for the pre-screening of quartz crystals for application in
     fluid property investigation. We also present a data fitting algorithm
     which enables the conversion of raw impedance data into equiv.
     circuit parameters. Subsequently, we report on our study of a
     series of Newtonian fluids (2-propanol/water solns.) by frequency response
     anal. of the fluid-contacted TSM QCR. The results are analyzed in
     comparison to theor. predictions presented in an earlier publication. The
     results show good agreement between the theory and exptl. derived equiv.
     circuit parameters. The influence of fluid elasticity on the
     impedance response of liq.-contacted thickness-shear mode (TSM) quartz
     crystal resonators (QCR) is investigated in the second part. Model
     predictions are compared to exptl. results on a series of a TSM QCR
     contacted with poly(dimethylsiloxane) fractions. The findings
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show that with appropriate instrumentation and models to interpret
     results, TSM QCR can be rapid and effective tools in viscoelastic fluid
    property investigation.
    crystal screening quartz resonator fluid property detn; thickness shear
ST
    mode quartz crystal resonator
ΙT
    Algorithm
        (data fitting; fluid property detn. by impedance of quartz crystal
       resonators)
IT
    Acoustic impedance
     Electric impedance
     Shear
    Thickness
     Viscoelastic materials
        (fluid property detn. by impedance of quartz crystal resonators)
ΙT
    Resonators
       (piezoelec.; fluid property detn. by impedance of quartz crystal
       resonators)
     14808-60-7, Quartz, uses
ΙT
     RL: DEV (Device component use); USES (Uses)
        (fluid property detn. by impedance of quartz crystal resonators)
     67-63-0, 2-Propanol, uses 7732-18-5, Water, uses
     RL: NUU (Other use, unclassified); USES (Uses)
        (fluid property detn. by impedance of quartz crystal resonators)
             THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT
RF.
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     14808-60-7
RN
     67-63-0
RN
RN
     7732-18-5
L10 ANSWER 16 OF 16 CAPLUS COPYRIGHT 2003 ACS
     2001:1253 CAPLUS
ΑN
     134:79701
DN
     Process for depositing a low dielectric constant film
TI
     Moghadam, Farhad; Cheung, David W.; Yieh, Ellie; Xia, Li-qun; Yau,
ΤN
     Wai-Fan; Lang, Chi-I.; Jeng, Shin-Puu; Gaillard, Frederick; Venkataraman,
     Shankar; Nemani, Srinivas D.
     Applied Materials, Inc., USA
PΑ
     Eur. Pat. Appl., 20 pp.
     CODEN: EPXXDW
DT
     Patent
    English
LA
     ICM H01L021-316
ΙC
     ICS C23C016-40
     76-3 (Electric Phenomena)
CC
     Section cross-reference(s): 75
FAN.CNT 8
                    KIND DATE
                                          APPLICATION NO. DATE
     PATENT NO.
     _____
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20001227
                      Α1
    EP 1063692
PΙ
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO
                                           US 1999-338470
                                                           19990622
                            20020702
     US 6413583
                      В1
                            19990622 5054379
PRAI US 1999-338470
                      Α
                           19980211 76 348723
19990210 76
                      A2
    US 1998-21788
    US 1999-247381
                     A2
     A method is presented for depositing Si oxide layers having a low dielec.
AΒ
     const. by reaction of an organosilicon compd. and a hydroxyl forming
     compd. at a substrate temp. .ltorsim.400.degree.. The low dielec. const.
     films contain residual C and are useful for gap fill layers, pre-metal
     dielec. layers, inter-metal dielec. layers, and shallow trench isolation
     dielec. layers in sub-micrometer devices. The hydroxyl compd. can be
     prepd. prior to deposition from H2O or an org. compd. The Si oxide layers
     are preferably deposited at a substrate temp. .ltorsim.40.degree. onto a
     liner layer produced from the organosilicon compd. to provide gap fill
     layers having a dielec. const. .ltorsim.3.0.
     plasma vapor deposition process dielec film
ST
     Semiconductor devices
ΙT
        (microscale, submicron; process for depositing a low dielec. const.
        film)
     Vapor deposition process
ΙT
        (plasma; process for depositing a low dielec. const. film)
     Dielectric films
TI
     Integrated circuits
     Semiconductor device fabrication
        (process for depositing a low dielec. const. film)
     7631-86-9, Silica, processes
ΙT
     RL: DEV (Device component use); PEP (Physical, engineering or chemical
     process); PROC (Process); USES (Uses)
        (process for depositing a low dielec. const. film)
     64-19-7, Acetic acid, processes 67-63-0, Isopropyl alcohol,
ΙT
                 7722-84-1, Hydrogen peroxide, processes 74087-85-7,
     processes
     Dimethyldioxirane
     RL: PEP (Physical, engineering or chemical process); RCT (Reactant); PROC
     (Process); RACT (Reactant or reagent)
        (vapor deposition hydroxyl-forming compd.; process for depositing a low
        dielec. const. film)
                                                               291-27-0,
                                  78-10-4, Tetraethoxysilane
     75-76-3, Tetramethylsilane
TΤ
     1,3,5-Trisilacyclohexane 293-36-7, 1,5-Dioxa-2,4,6,8-
     tetrasilacyclooctane 992-94-9, Methylsilane 993-07-7, Trimethylsilane
     1111-74-6, Dimethylsilane 1759-88-2 3277-26-7, 1,1,3,3-
                            4364-07-2 4405-22-5
                                                     4745-36-2
     Tetramethyldisiloxane
                6166-86-5, 2,4,6,8,10-Pentamethylcyclopentasiloxane
     5654-05-7
                 78570-62-4 234772-32-8
                                            234772-34-0
     14396-21-5
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (vapor deposition precursor; process for depositing a low dielec.
        const. film)
              THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT 5
RE
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     7631-86-9
RN
RN
     64-19-7
     67-63-0
RN
     7722-84-1
RN
RN
     74087-85-7
RN
     75-76-3
RN
     78-10-4
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EP 2000-112820

RN291-27-0 293-36-7 RN 992-94-9 RN993-07-7 RN 1111-74-6 RN 1759-88-2 RN 3277-26-7 RN4364-07-2 RN4405-22-5 RN 4745-36-2 RN5654-05-7 RN6166-86-5 RN14396-21-5 RN78570-62-4 RN 234772-32-8 RN 234772-34-0 RN

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L13 ANSWER 1 OF 15 CAPLUS COPYRIGHT 2003 ACS
    2000:419060 CAPLUS
AN
    133:273888
DN
    Covalent bonding of coumarin molecules to sol-gel matrices for organic
ΤI
     light-emitting device applications
     Karkkainen, Ari H. O.; Hormi, Osmo E. O.; Rantala, Juha T.
ΑU
    VTT Electronics, Oulu, Finland
CS
     Proceedings of SPIE-The International Society for Optical Engineering
SO
     (2000), 3943(Sol-Gel Optics V), 194-209
     CODEN: PSISDG; ISSN: 0277-786X
     SPIE-The International Society for Optical Engineering
PΒ
DT
     Journal
     English
LΑ
     73-11 (Optical, Electron, and Mass Spectroscopy and Other Related
CC
     Properties)
     Section cross-reference(s): 27, 41, 57
     Coumarin mols. are widely used as laser dyes and their luminescence
AΒ
     properties show a large potential for their use as light emitters in org.
     light emitting devices. These mols. however are lacking of photo, chem.
     and thermal stability. At the outset, the fact that when the coumarin or
     other org. active mols. are covalently bonded to a metal oxide host, the
     stability properties can be improved. The authors outline the synthesis
     of several different coumarin-3-carboxylic acids by using a 1-pot
     synthesis from dipotassium o-methoxybenzylidenemalonates. The authors
     also outline a preparative route for the synthesis of corresponding
     coumarin-3-carboxylic amides with a side chain contg. terminal
     trimethyoxysilane functionality, which allows the creation of a covalent
     bond between the mol. and a Si oxide host matrix. These silylated
     coumarins are then covalently bonded through a sol-gel method to a
     developing siloxane host matrix. The Si matrix materials were
     synthesized through hydrolysis and simultaneous condensation of
     metalalkoxides such as phenylmethyltrimethoxysilane. Coumarin dyes are
     bonded in- situ to the developing matrix during the prepn. of the matrix.
     The excitation and emission spectra of these mols. are examd. in liq.
     phase to evaluate the effect of varying substitution pattern on
     luminescence characteristics. The photo luminescence characteristics are
     also measured from a solid thin film to explore the effect of the matrix
     on emission wavelengths. These materials show potentiality for their
     applications in thin film electro luminescence devices whose fabrication
     and properties are finally discussed.
     coumarin dye deriv covalent bond siloxane sol gel LED; LUMO
ST
     coumarin dye deriv hybrid siloxane sol gel LED; HOMO
     coumarin dye deriv hybrid siloxane sol gel LED;
     structure luminescence coumarin dye deriv hybrid
     siloxane sol gel; substituent coumarin dye deriv hybrid
     siloxane sol gel luminescence; mass spectra coumarin dye deriv
     hybrid siloxane sol gel; NMR coumarin dye deriv
     hybrid siloxane sol gel; melting point coumarin dye
     deriv hybrid siloxane sol gel; amide coumarin dye
     deriv hybrid siloxane sol gel LED; carboxylic acid
     coumarin dye deriv siloxane sol gel LED
     Carboxylic acids, properties
ΙT
     RL: DEV (Device component use); PRP (Properties); SPN (Synthetic
     preparation); PREP (Preparation); USES (Uses)
        (coumarin-3-; covalent bonding of coumarin mols. to sol-gel matrixes
        for org. light-emitting device applications)
     Amides, properties
IT
     RL: DEV (Device component use); PRP (Properties); SPN (Synthetic
     preparation); PREP (Preparation); USES (Uses)
        (coumarin-3-carboxylic; covalent bonding of coumarin mols. to sol-gel
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matrixes for org. light-emitting device applications)
    Bond formation
TΤ
    Differential scanning calorimetry
    Electroluminescent devices
    HOMO (molecular orbital)
    LUMO (molecular orbital)
     Luminescence
    Mass spectra
    Melting point
    NMR (nuclear magnetic resonance)
     Sol-gel processing
     Substituent effects
        (covalent bonding of coumarin mols. to sol-gel matrixes for org.
        light-emitting device applications)
     Siloxanes (nonpolymeric)
IT
     RL: DEV (Device component use); PEP (Physical, engineering or chemical
     process); RCT (Reactant); PROC (Process); RACT (Reactant or reagent); USES
     (Uses)
        (covalent bonding of coumarin mols. to sol-gel matrixes for org.
        light-emitting device applications)
ΙT
     Dyes
        (laser, coumarin; covalent bonding of coumarin mols. to sol-gel
        matrixes for org. light-emitting device applications)
     Molecular structure-property relationship
IT
        (luminescence; covalent bonding of coumarin mols. to sol-gel matrixes
        for org. light-emitting device applications)
                                110-89-4, Piperidine, uses
     64-19-7, Acetic acid, uses
ΙT
     RL: CAT (Catalyst use); USES (Uses)
        (covalent bonding of coumarin mols. to sol-gel matrixes for org.
        light-emitting device applications)
TΥ
     2996-92-1
     RL: DEV (Device component use); PEP (Physical, engineering or chemical
     process); PRP (Properties); RCT (Reactant); PROC (Process); RACT (Reactant
     or reagent); USES (Uses)
        (covalent bonding of coumarin mols. to sol-gel matrixes for org.
        light-emitting device applications)
                                            2085-33-8, Hydroxyquinoline
     531-81-7, Coumarin-3-carboxylic acid
IΤ
     aluminum
     RL: PRP (Properties)
        (covalent bonding of coumarin mols. to sol-gel matrixes for org.
        light-emitting device applications)
                                            4460-86-0P, 2,4,5-
     613-45-6P, 2,4-Dimethoxybenzaldehyde
ΙT
                             7324-86-9P, Diethyl-2,4-
     Trimethoxybenzaldehyde
                                   57724-31-9P, Diethyl-2,4,5-
     dimethoxybenzylidenemalonate
     trimethoxybenzylidenemalonate 84562-48-1P, 4-Dimethylamino-2-
                                        133031-78-4P, Dipotassium-2,4-
                           86726-44-5P
     methoxybenzaldehyde
     dimethoxybenzylidenemalonate 133031-81-9P, Dipotassium-2,4,5-
                                                                   297736-08-4P
                                                   297736-07-3P
                                    297736-06-2P
     trimethoxybenzylidenemalonate
     RL: PRP (Properties); RCT (Reactant); SPN (Synthetic preparation); PREP
     (Preparation); RACT (Reactant or reagent)
        (covalent bonding of coumarin mols. to sol-gel matrixes for org.
        light-emitting device applications)
     20300-59-8P, 7-Methoxycoumarin-3-carboxylic acid 55804-65-4P, Coumarin
ΙT
                         86100-68-7P, 6,7-Dimethoxycoumarin-3-carboxylic acid
           81017-27-8P
                                                  297736-11-9P 297736-12-0P
                                  297736-10-8P
     122607-15-2P
                    297736-09-5P
     297736-13-1P
     RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
         (covalent bonding of coumarin mols. to sol-gel matrixes for org.
        light-emitting device applications)
                                      76-05-1, Trifluoroacetic acid,
     67-63-0, 2-Propanol, reactions
 IT
                                             407-25-0, Trifluoroacetic
                 105-53-3, Diethyl malonate
      reactions
                 530-62-1, 1,1'-Carbonyldiimidazole 830-79-5,
      anhydride
```

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1310-58-3, Potassium hydroxide, reactions
    2,4,6-Trimethoxybenzaldehyde
    13822-56-5
    RL: RCT (Reactant); RACT (Reactant or reagent)
        (covalent bonding of coumarin mols. to sol-gel matrixes for org.
        light-emitting device applications)
              THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD
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RN
     110-89-4
RN
RN
     2996-92-1
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     531-81-7
    2085-33-8
RN
    613-45-6P
RN
    4460-86-0P
RN
    7324-86-9P
RN
    57724-31-9P
RN
    84562-48-1P
RN
    86726-44-5P
RN
RN
    133031-78-4P
    133031-81-9P
RN
     297736-06-2P
RN
    297736-07-3P
RN
    297736-08-4P
RN
     20300-59-8P
RN
     55804-65-4P
RN
     81017-27-8P
RN
RN
     86100-68-7P
RN
     122607-15-2P
RN
     297736-09-5P
RN
     297736-10-8P
     297736-11-9P
RN
RN
     297736-12-0P
RN
     297736-13-1P
RN
     67-63-0
RN
     76-05-1
RN
     105-53-3
RN
     407-25-0
     530-62-1
RN
     830-79-5
RN
     1310-58-3
RN
     13822-56-5
RN
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L13 ANSWER 2 OF 15 CAPLUS COPYRIGHT 2003 ACS

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2002:307198 CAPLUS
AΝ
     137:116871
DN
     Siloxane-based hybrid glass materials for binary and
TТ
     gray-scale mask photoimaging
    Karkkainen, Ari H. O.; Rantala, Juha T.; Maaninen, Arto; Jabbour, Ghassan
ΑU
     E.; Descour, Michael R.
     VTT Electronics, Oulu, FIN-90570, Finland
CS
     Advanced Materials (Weinheim, Germany) (2002), 14(7), 535-540
SO
     CODEN: ADVMEW; ISSN: 0935-9648
     Wiley-VCH Verlag GmbH
PΒ
     Journal
DT
     English
LΑ
     74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other
CC
     Reprographic Processes)
     Section cross-reference(s): 73
     The fabrication of microoptical and optomech. structures by applying
AΒ
     photoimaging of hybrid glass materials is discussed. The
     optical and optomech. structures are fabricated simultaneously in a single
     lithog. step. Gray-scale and binary photomasks have been successfully
     applied for the fabrication of lens arrays to a max. lens sag of 102 .mu.m
     and of optomech. structures to a max. height of 140 .mu.m.
     Alignment-aiding optomech. structures can be patterned simultaneously with
     optical structures in the hybrid glass to fabricate microoptical
     elements. No chem. or dry etch transfer of the imaged structures is
     required. The fabricated lenslets and the optomech. structures show high
     surface and optical quality. The fabricated hybrid glass
     surfaces can be coated with interference coatings utilizing std.
     deposition procedures. Photoimaging of hybrid glass materials
     simplifies the fabrication of the optical components and enables new
     optics integration options.
     hybrid glass siloxane based photoimaging microoptical
ST
     optomech structure; photolithog hybrid glass siloxane
     based microoptical optomech structure fabrication
     Photoimaging materials
ΙT
     Photolithography
     Surface roughness
        (photoimaging of siloxane-based neg-tone hybrid
        glass materials in fabrication of microoptical and optomech.
        structures)
     Polysiloxanes, processes
ΙT
     RL: PEP (Physical, engineering or chemical process); PYP (Physical
     process); PROC (Process)
        (photoimaging of siloxane-based neg-tone hybrid
        glass materials in fabrication of microoptical and optomech.
        structures)
     Ceramers
ΤТ
       Hybrid organic-inorganic materials
        (photoimaging of siloxane-based neg-tone hybrid
        glass materials in fabrication of microoptical and optomech. structures
        in relation to)
     Polymerization
IΤ
        (photopolymn.; photoimaging of siloxane-based neg-tone
        hybrid glass materials in fabrication of microoptical and
        optomech. structures)
     Microlenses
IT
        (refractive; photoimaging of siloxane-based neg-tone
        hybrid glass materials in fabrication of microoptical and
        optomech. structures)
     442874-00-2P, Phenyltrimethoxysilane-[3-(Methacryloyloxy)propyl]trimethoxy
ΙT
     silane-trimethylolpropane trimethacrylate copolymer
     RL: PEP (Physical, engineering or chemical process); PYP (Physical
     process); SPN (Synthetic preparation); PREP (Preparation); PROC (Process)
```

(crosslinked; photoimaging of siloxane-based neg-tone hybrid glass materials in fabrication of microoptical and optomech. structures) 108-10-1, Methyl isobutyl ketone 67-63-0, Isopropanol, uses TΥ RL: NUU (Other use, unclassified); USES (Uses) (developer mixt.; photoimaging of siloxane-based neg-tone hybrid glass materials in fabrication of microoptical and optomech. structures) 947-19-3, 1-Hydroxycyclohexylphenyl ketone 162881-26-7 TΤ RL: CAT (Catalyst use); USES (Uses) (photopolymn. initiator system; photoimaging of siloxane -based neg-tone hybrid glass materials in fabrication of microoptical and optomech. structures) IT 94-36-0, Benzoyl peroxide, uses RL: CAT (Catalyst use); USES (Uses) (prepolymer soln.; photoimaging of siloxane-based neg-tone hybrid glass materials in fabrication of microoptical and optomech. structures) 2996-92-1, 2530-85-0, 3-[Methacryloyloxy)propyl]trimethoxysilane TΨ Phenyltrimethoxysilane 3290-92-4, Trimethylolpropane trimethacrylate RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PYP (Physical process); RCT (Reactant); PROC (Process); RACT (Reactant or reagent) (prepolymer soln.; photoimaging of siloxane-based neg-tone hybrid glass materials in fabrication of microoptical and optomech. structures) THERE ARE 27 CITED REFERENCES AVAILABLE FOR THIS RECORD RE.CNT (1) Ayraes, P; Thin Solid Films 1999, V352, P9 (2) Conrady, A; Applied Optics and Optical Design Part 2 1960 (3) Cox, W; Opt Photon News 2001, V12(6), P32 CAPLUS (4) Dannberg, P; Microsyst Technol 1999, V6, P41 (5) Decker, C; Makromol Chem 1988, V189, P2381 CAPLUS (6) Eisner, M; Opt Eng 1996, V10, P2979 (7) Fouassier, J; J Photochem Photobiol A: Chem 1991, V61, P47 CAPLUS (8) Fouassier, J; Photoinitiation, Photopolymerization and Photocuring: Fundamentals and Applications 1995 (9) Gerlach, A; Microsyst Technol 2001, V7, P27 (10) Haruvy, Y; Chem Mater 1991, V3, P501 CAPLUS (11) Haruvy, Y; Chem Mater 1997, V9, P2604 CAPLUS (12) Keyworth, B; US 5534101 1996 (13) Krug, H; New J Chem 1994, V18, P1125 CAPLUS (14) Kupka, R; Appl Surf Sci 2000, V164, P97 CAPLUS (15) Levy, R; Proc SPIE-Int Soc Opt Eng 1999, V3879, P167 (16) Mohr, J; Sens Mater 1998, V10, P363 CAPLUS (17) Morris, G; Proc SPIE-Int Soc Opt Eng 1998, V3573, P370 CAPLUS (18) Noll, R; J Opt Soc Am 1976, V66, P207 (19) Rantala, J; Electron Lett 2000, V16, P530 (20) Rantala, J; Opt Lett 1998, V23, P1939 CAPLUS (21) Riedl, M; Photon Spectra 2001, V35, P130 (22) Schaffer, C; Opt Photon News 2001, V12(4), P20 (23) Seraji, S; Adv Mater 2000, V12, P1421 CAPLUS (24) Shannon, R; The Art and Science of Optical Design 1997 (25) Shinmou, K; J Sol-Gel Sci Technol 2000, V19, P267 CAPLUS (26) Sinzinger, S; Microoptics 1999 (27) Yoldas, B; J Sol-Gel Sci Technol 1998, V13, P147 CAPLUS RN 442874-00-2P RN 67-63-0 RN 108-10-1 RN 947-19-3 RN 162881-26-7

RN

94-36-0

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RN
    2530-85-0
RN
    2996-92-1
RN 3290-92-4
L13 ANSWER 3 OF 15 CAPLUS COPYRIGHT 2003 ACS
   2002:483594 CAPLUS
DN
    137:144003
     Preparation of inorganic-organic hybrid films containing
TΙ
     particles using electrophoretic deposition method
     Yamada, Noriko; Shoji, Hiromasa; Kubo, Yuji; Katayama, Shingo
ΑIJ
    Advanced Technology Research Laboratories, Nippon Steel Corporation,
CS
     Futtsu, Chiba, 293-8511, Japan
     Journal of Materials Science (2002), 37(10), 2071-2076
SO
     CODEN: JMTSAS; ISSN: 0022-2461
PB
    Kluwer Academic Publishers
    Journal
DT
     English
LA
     57-1 (Ceramics)
CC
     Section cross-reference(s): 55
     The composite films of methylsiloxane inorg.-org. hybrid
AΒ
     and MoS2 particles have successfully been fabricated by electrophoretic
     deposition of MoS2 particles in a mixed soln. of Me Et ketone (MEK) and
     inorg.-org. hybrid sol. The addn. of 20 vol% hybrid
     sol into a MEK suspension increased the amt. of MoS2 deposition twice as
     much as that of MoS2 deposition in MEK alone. The fraction of particles
     deposited on a substrate in MEK-20 vol* sol was estd. to be much larger
     than that in MEK. The hydrolyzed methyltriethoxysilane in a
     hybrid sol modified the MoS2 particles, resulting in lower neg.
     zeta potential, which reduces the repulsion force among particles and
     makes the incorporation of particles into a deposition film easier. The
     surface modification also enables the incorporation of particles into a
     deposit by the interaction of surface modifiers. These factors enhance
     the incorporation of MoS2 particles in electrophoretic deposition in
     inorg org hybrid molybdenum sulfide composite film
ST
     electrophoretic deposition; silica gel molybdenum sulfide composite film
     electrophoretic deposition
     Electrophoretic deposition
ΙT
     Zeta potential
        (electrophoretic deposition of organically modified silica-MoS2
        particle composite films)
     Silica gel, preparation
ΙT
     RL: PEP (Physical, engineering or chemical process); PRP (Properties); PYP
     (Physical process); SPN (Synthetic preparation); PREP (Preparation); PROC
     (Process)
        (organically modified, composite films; electrophoretic deposition of
        organically modified silica-MoS2 particle composite films)
ΤТ
     Ceramers
       Hybrid organic-inorganic materials
        (silica-molybdenum sulfide composite films; electrophoretic deposition
        of organically modified silica-MoS2 particle composite films)
     1317-33-5, Molybdenum sulfide (MoS2), processes
TΤ
     RL: PEP (Physical, engineering or chemical process); PRP (Properties); PYP
     (Physical process); PROC (Process)
        (composite films; electrophoretic deposition of organically modified
        silica-MoS2 particle composite films)
ΙT
     60676-86-0P, Vitreous silica
     RL: PEP (Physical, engineering or chemical process); PRP (Properties); PYP
     (Physical process); SPN (Synthetic preparation); PREP (Preparation); PROC
     (Process)
        (gel-glass, organically modified, composite films; electrophoretic
```

deposition of organically modified silica-MoS2 particle composite

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films)
    2031-67-6, Silane, triethoxymethyl-
IT
    RL: CPS (Chemical process); PEP (Physical, engineering or chemical
    process); PROC (Process)
        (precursor; electrophoretic deposition of organically modified
       silica-MoS2 particle composite films)
    64-17-5, Ethanol, uses 67-63-0, 2-Propanol, uses
ΙT
    Acetone, uses 78-93-3, Methyl ethyl ketone, uses 108-10-1, Methyl
    isobutyl ketone 110-80-5, 2-Ethoxyethanol 123-54-6, Acetylacetone,
           141-97-9, Ethyl acetoacetate
    RL: MOA (Modifier or additive use); USES (Uses)
       (solvent; electrophoretic deposition of organically modified
       silica-MoS2 particle composite films)
ΙT
    11109-50-5, Sus 304
    RL: NUU (Other use, unclassified); USES (Uses)
       (substrates; electrophoretic deposition of organically modified
       silica-MoS2 particle composite films)
             THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD
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   1317-33-5
RN
    60676-86-0P
RN
    2031-67-6
RN
   64-17-5
RN
RN
    67-63-0
RN
    67-64-1
    78-93-3
RN
RN
    108-10-1
RN
    110-80-5
RN
    123-54-6
RN
    141-97-9
    11109-50-5
RN
L13 ANSWER 4 OF 15 CAPLUS COPYRIGHT 2003 ACS
    1998:204034 CAPLUS
NΑ
    129:33730
DN
    Electrochemical impedance studies of hybrids of
    solution
     Zoppi, R. A.; Nunes, S. P.
ΑU
    Instituto de Quimica, Universidade Estadual de Campinas, CEP 13083-970,
CS
     Campinas, Brazil
    Journal of Electroanalytical Chemistry (1998), 445(1-2), 39-45
SO
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CODEN: JECHES; ISSN: 0368-1874
PΒ
    Elsevier Science S.A.
DT
    Journal
LA
    English
    72-3 (Electrochemistry)
CC
     Section cross-reference(s): 37, 76
     Hybrids of Nafion and silica were prepd. from soln., growing the
AB
     inorg. phase by hydrolysis/condensation of alkoxy silanes. Using
     tetraethoxysilane (TEOS) as the inorg. precursor, transparent and rigid
     films were obtained. Substituting part of the TEOS (20 wt substitution
     by 1,1,3,3- tetramethyl-1,3-diethoxydisiloxane (TMDES) more
     flexible films were obtained. These films were translucent and showed a
     phase segregation which was clearly obsd. by transmission electron
     microscopy. The ionic cond. of the hybrids was measured by
     electrochem. impedance spectroscopy using two stainless steel electrodes,
     a frequency range of 0.1 to 105 Hz, and temps. from 25 to 100.degree.C.
     Samples were also characterized by modulated differential scanning
     calorimetry.
     hybrid material composite ionic conductor Nafion; silica nafion
ST
     hybrid hydrolysis condensation alkoxysilanes
ΙT
     Spectroscopy
        (energy-dispersive; of matrix and domain in Nafion/TEOS/TMDS
        hvbrids)
IT
     Polyoxyalkylenes, properties
     RL: PEP (Physical, engineering or chemical process); PRP (Properties); RCT
     (Reactant); PROC (Process); RACT (Reactant or reagent)
        (fluorine- and sulfo-contg., ionomers; hybrids with silicon
        oxide by sol-gel reaction from soln.)
     Polyoxyalkylenes, properties
ΙT
     RL: PEP (Physical, engineering or chemical process); PRP (Properties); RCT
     (Reactant); PROC (Process); RACT (Reactant or reagent)
        (fluorine-contg., sulfo-contg., ionomers; hybrids with
        silicon oxide by sol-gel reaction from soln.)
     Ionic conductivity
ΙT
        (for Nafion/TEOS/TMDES systems)
ΤТ
     Activation energy
        (for distribution of free vol. in Nafion/TEOS/TMDES hybrids)
     Differential scanning calorimetry
IT
        (for pure Nafion)
TΤ
     Ionomers
     RL: PEP (Physical, engineering or chemical process); PRP (Properties); RCT
     (Reactant); PROC (Process); RACT (Reactant or reagent)
        (hybrids with silicon oxide by sol-gel reaction from soln.)
ΙT
     Phase separation
        (in films of hybrids of Nafion/TEOS/TMDES by TEM)
     Condensation reaction
ΙT
     Hydrolysis
        (of alkoxysilanes in formation of hybrids of
        Nafion/TEOS/TMDES)
     Electric impedance
ΙT
        (of hybrids films of silica with perfluorosulfonic acid
        ionomer)
ΙT
     Films
        (of hybrids of Nafion/TEOS/TMDES; formation and properties)
IΤ
     Surface structure
        (of pure Nafion films and Nafion/TEOS/TMDES hybrids by TEM)
     Fluoropolymers, properties
ΙT
     Fluoropolymers, properties
     RL: PEP (Physical, engineering or chemical process); PRP (Properties); RCT
     (Reactant); PROC (Process); RACT (Reactant or reagent)
        (polyoxyalkylene-, sulfo-contg., ionomers; hybrids with
        silicon oxide by sol-gel reaction from soln.)
```

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TT
    Ionomers
    RL: PEP (Physical, engineering or chemical process); PRP (Properties); RCT
     (Reactant); PROC (Process); RACT (Reactant or reagent)
        (polyoxyalkylenes, fluorine- and sulfo-contg.; hybrids with
        silicon oxide by sol-gel reaction from soln.)
ΙT
     Phase transition
        (reversible, exothermic: related to cluster reorganization)
IΤ
    7631-86-9, Silica, properties
     RL: PEP (Physical, engineering or chemical process); PRP (Properties); RCT
     (Reactant); PROC (Process); RACT (Reactant or reagent)
        (hybrids with perfluorosulfonic acid ionomer by sol-gel
        reaction from soln.)
ΙT
     7732-18-5, Water, uses
     RL: NUU (Other use, unclassified); PRP (Properties); USES (Uses)
        (in fabrication of hybrids of silica with perfluorosulfonic
        acid ionomer by sol-gel reaction from soln.)
                                         71-23-8, n-Propanol,
     67-63-0, Iso-propanol, properties
TΤ
                 7647-01-0, Hydrochloric acid, properties
    properties
     RL: PEP (Physical, engineering or chemical process); PRP (Properties); RCT
     (Reactant); PROC (Process); RACT (Reactant or reagent)
        (in fabrication of hybrids of silica with perfluorosulfonic
        acid ionomer by sol-gel reaction from soln.)
ΙΤ
     18420-09-2, 1,1,3,3-Tetramethyl-1,3-diethoxydisiloxane
     RL: MOA (Modifier or additive use); PEP (Physical, engineering or chemical
     process); PRP (Properties); RCT (Reactant); PROC (Process); RACT (Reactant
     or reagent); USES (Uses)
        (precursor for SiO2 obtaining in formation of hybrids of
        Nafion and silica)
    78-10-4, Tetraethoxysilane
ΤТ
     RL: PEP (Physical, engineering or chemical process); PRP (Properties); RCT
     (Reactant); PROC (Process); RACT (Reactant or reagent)
        (precursor for SiO2 obtaining in formation of hybrids of
        Nafion and silica)
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     7631-86-9
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     7732-18-5
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RN
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     7647-01-0
RN
     18420-09-2
RN
RN
     78-10-4
L13 ANSWER 5 OF 15 CAPLUS COPYRIGHT 2003 ACS
AN
     1989:81194 CAPLUS
DN
     110:81194
     Solvent extraction of inorganic/organic hybrid gels prepared by
ΤI
     the sol-gel method
ΑU
     Kojiya, Shinzo; Ochiai, Kenichiro; Yamashita, Shinzo
CS
     Dep. Chem., Kyoto Inst. Technol., Kyoto, 606, Japan
     Chemistry Express (1988), 3(10), 631-4
SO
     CODEN: CHEXEU; ISSN: 0911-9566
DT
     Journal
LA
     Japanese
CC
     57-9 (Ceramics)
     Solvent extns. were carried out on inorg./org. hybrid gels
AΒ
     prepd. by the sol-gel method. Incorporation of hydroxy-terminated poly(
     dimethylsiloxane) into glass matrix was nonquant. About 20 - of
     the original polymer remained unreacted and could be extd. TG anal. of
     the gels indicated that the incorporated polymer has better thermal
     stability than the free polymer.
     inorg org hybrid sol gel; solvent extn inorg org gel
ST
     hybrid
ΙT
     Gels
        (hybrid, inorg.-org., solvent extn. from)
TΤ
     Polymers, uses and miscellaneous
     RL: USES (Uses)
        (inorg. gel hybrid with, solvent extn. from)
IΤ
     67-63-0, 2-Propanol, uses and miscellaneous 75-09-2,
     Dichloromethane, uses and miscellaneous 108-88-3, Toluene, uses and
     miscellaneous 109-99-9, THF, uses and miscellaneous
     Cyclohexane, uses and miscellaneous 123-91-1, Dioxane, uses and
     miscellaneous
     RL: USES (Uses)
        (extn. of, from inorg.-org. hybrid gels)
RN
     67-63-0
RN
     75-09-2
RN
     108-88-3
RN
     109-99-9
RN
     110-82-7
RN
     123-91-1
L13 ANSWER 6 OF 15 CAPLUS COPYRIGHT 2003 ACS
     2001:737029 CAPLUS
AN
DN
     135:305276
ΤŢ
     Primer for use under a photocatalytic coating layer and coated articles
     using the primer
ΙN
     Nakanishi, Makoto; Kojima, Eiichi; Kanamori, Taro; Hashiguchi, Yuichi
    Toto Ltd., Japan; JSR Ltd.
PA
     Jpn. Kokai Tokkyo Koho, 10 pp.
     CODEN: JKXXAF
DT
    Patent
LA
     Japanese
TC
     ICM C09D201-00
     ICS B01J035-02; B01J037-02; B32B009-00; C09D005-00; C09D143-00;
          C09D183-04
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42-10 (Coatings, Inks, and Related Products)
FAN.CNT 1
                                         APPLICATION NO. DATE
    PATENT NO.
                    KIND DATE
                                         -----
    _____
                                        JP 2000-97952 20000331
    JP 2001279184 A2 20011010
PΙ
PRAI JP 2000-97952
                          20000331
   The primer can prevent the erosion of coated articles such as exterior
    plastics due to the photolytic attack, and contains hybrid
    org.-inorg. materials and solvents which have SP value difference with
    that of the substrate of 0.2-1.6. Thus, mixing PMMA 36 with a
    polysiloxane 24, i-PrOH (SP value 11.50) 285, BuOAc (SP value
     8.50) 285, MEK (SP value 9.30) 285 and EtOAc (SP value 9.10) 285 parts 🕢
     gave a primer which was coated on the cleaned surface of a Sumipex Clear
     000 (PMMA; SP value 9.10) panel to a wet pickup wt. of 20-25 g/m2, dried
     at 80.degree. for 30 min with hot air, cooled to room temp., over-coared
     with a photocatalytic coating contg. ST-K 01 and ST-K 03 and dried t\phi' give
     a panel with lasting wettability and coat film adhesion.
    photocatalytic coating protection primer hybrid org inorg
ST
    material
    Photolysis catalysts
ΤT
        (coatings; primer for use under a photolytic coating layer and coated
        articles)
    Polysiloxanes, uses
ΙT
     RL: POF (Polymer in formulation); RCT (Reactant); TEM (Technical or
     engineered material use); RACT (Reactant or reagent); USES (Uses)
        (hybrid org.-inorg. materials for primer; primer for use
        under a photolytic coating layer and coated articles)
IT
        (org.; primer for use under a photolytic coating layer and coated
        articles)
     Hybrid organic-inorganic materials
ΤТ
     Primers (paints)
     Solvent effect
        (primer for use under a photolytic coating layer and coated articles)
ΙT
     9011-14-7, PMMA
     RL: POF (Polymer in formulation); RCT (Reactant); TEM (Technical or
     engineered material use); RACT (Reactant or reagent); USES (Uses)
        (hybrid org.-inorg. materials for primer; primer for use
        under a photolytic coating layer and coated articles)
     202936-29-6, ST-K 03 220946-52-1, ST-K 01
IΤ
     RL: TEM (Technical or engineered material use); USES (Uses)
        (photocatalytic coatings; primer for use under a photolytic coating
        layer and coated articles)
     67-63-0, Isopropanol, uses
                                78-93-3, MEK, uses 123-86-4, Butyl
ΤТ
     acetate 141-78-6, Ethyl acetate, uses
     RL: NUU (Other use, unclassified); USES (Uses)
        (solvents of specified soly. parameter; primer for use under a
        photolytic coating layer and coated articles)
     9011-14-7
RN
     202936-29-6
RN
    220946-52-1
RN
     67-63-0
RN
     78-93-3
RN
     123-86-4
RN
    141-78-6
RN
L13 ANSWER 7 OF 15 CAPLUS COPYRIGHT 2003 ACS
     1999:390437 CAPLUS
ΑN
     131:40530
DN
    Method for identifying nucleic acids by means of matrix-assisted laser
TΙ
     desorption/ionisation mass spectrometry using charge tag-labeled
     hybridization probes
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Max-Planck-Gesellschaft zur Forderung der Wissenschaften e.V., Germany
PΑ
SO
    PCT Int. Appl., 51 pp.
    CODEN: PIXXD2
DT
    Patent
LΑ
    German
    ICM C12Q001-68
TC
    3-1 (Biochemical Genetics)
    Section cross-reference(s): 7, 9
FAN.CNT 1
                                        APPLICATION NO. DATE
                    KIND DATE
    PATENT NO.
                                         _____
                    ----
                    A2 19990617
                                        WO 1998-EP7911 19981204
    WO 9929898
PΤ
    WO 9929898
                    A3 19991028
        W: CA, JP, US
        RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL,
            PT, SE
                         19990617
                                         CA 1998-2312052 19981204
                      AA
    CA 2312052
                     A2 20000920
                                         EP 1998-966608
                                                          19981204
    EP 1036202
                    B1 20020502
    EP 1036202
        R: AT, BE, CH, DE, DK, ES, FR, GB, IT, LI, NL, SE, PT, IE, FI
    JP 2001526381 T2 20011218 JP 2000-524469
                                                          1998120/4
                     E
                          20020515
                                         AT 1998-966608
    AT 217028
                     T3 20021016
                                         ES 1998-966608
                                                          19981204
    ES 2173670
                   A 19971205
W 19981204
PRAI EP 1997-121471
    WO 1998-EP7911
    A method for detecting a nucleotide sequence in a nucleic a\phiid mol. by
AB
    means of hybridization against an ordered array of charged /
     tag-labeled probes using MALDI mass spectrometry is described.
     advantage of the method is that it provides for simultaneous
     characterization of a no. of unknown nucleic acid mols. using this array
     of probes. In particular peptide analogs of nucleic acids, esp. peptide
     nucleic acids, that showed improved efficiency of desorption are used as
     probes. The invention also relates to a kit contg. the probes and/or a
     sample carrier, optionally with nucleic acids bonded thereto.
    MALDI immobilized probe DNA sequencing
ST
    Proteins, specific or class
ΤТ
     RL: DEV (Device component use); MOA (Modifier or additive use); RCT
     (Reactant); RACT (Reactant or reagent); USES (Uses)
        (gene 32, in immobilization of hybridization of probes on
       MALDI matrixes; method for identifying nucleic acids by means of
       matrix-assisted laser desorption/ionisation mass spectrometry using
       charge tag-labeled hybridization probes)
     Peptide nucleic acids
IT
     Phosphorothioate oligonucleotides
     RL: ARG (Analytical reagent use); ANST (Analytical study); USES (Uses)
        (immobilized, as desorbable probes; method for identifying nucleic
       acids by means of matrix-assisted laser desorption/ionisation mass
       spectrometry using charge tag-labeled hybridization probes)
TΤ
    Amines, reactions
     Antibodies
     Epoxides
      Polysiloxanes, reactions
     Silanes
     Thiols (organic), reactions
     RL: DEV (Device component use); MOA (Modifier or additive use); RCT
     (Reactant); RACT (Reactant or reagent); USES (Uses)
        (in immobilization of hybridization of probes on MALDI
       matrixes; method for identifying nucleic acids by means of
       matrix-assisted laser desorption/ionisation mass spectrometry using
        charge tag-labeled hybridization probes)
     DNA sequence analysis
ΙT
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Gut, Ivo Glynne; Berlin, Kurt; Lehrach, Hans

ΙN

```
(mass spectrometric; method for identifying nucleic acids by means of
       matrix-assisted laser desorption/ionisation mass spectrometry using
       charge tag-labeled hybridization probes)
    Nucleic acid hybridization
TΤ
        (method for identifying nucleic acids by means of matrix-assisted laser
       desorption/ionisation mass spectrometry using charge tag-labeled
       hybridization probes)
IΤ
     Combinatorial library
        (of charge tag labeled probes; method for identifying nucleic acids by
       means of matrix-assisted laser desorption/ionisation mass spectrometry
       using charge tag-labeled hybridization probes)
ΙT
     Immobilization, biochemical
        (of probes on matrixes for MALDI; method for identifying nucleic acids
       by means of matrix-assisted laser desorption/ionisation mass
        spectrometry using charge tag-labeled hybridization probes)
     Probes (nucleic acid)
IT
     RL: ARG (Analytical reagent use); ANST (Analytical study); USES (Uses)
        (ordered arrays, charge labeled; method for identifying nucleic acids
       by means of matrix-assisted laser desorption/ionisation mass
        spectrometry using charge tag-labeled hybridization probes)
     Laser ionization mass spectrometry
ΙT
        (photodesorption, matrix-assisted; method for identifying nucleic acids
       by means of matrix-assisted laser desorption/ionisation mass
        spectrometry using charge tag-labeled hybridization probes)
     Laser desorption mass spectrometry
ΤT
        (photoionization, matrix-assisted; method for identifying nucleic acids
       by means of matrix-assisted laser desorption/ionisation mass
        spectrometry using charge tag-labeled hybridization probes)
     1519-55-7, .alpha.-Cyano-4-methoxycinnamic acid 227080-33-3
ΙT
     227080-34-4
     RL: DEV (Device component use); USES (Uses)
        (as matrix for MALDI; method for identifying nucleic acids by means of
       matrix-assisted laser desorption/ionisation mass spectrometry using
        charge tag-labeled hybridization probes)
     64-17-5, Ethanol, uses 67-56-1, Methanol, uses 67-63-0,
ΙT
                      67-64-1, 2-Propanone, uses 75-05-8, Acetonitrile,
     2-Propanol, uses
           7732-18-5, Water, uses
     uses
     RL: TEM (Technical or engineered material use); USES (Uses)
        (as solvent; method for identifying nucleic acids by means of
       matrix-assisted laser desorption/ionisation mass spectrometry using
        charge tag-labeled hybridization probes)
     58-85-5, Biotin 9013-20-1, Streptavidin
ΙT
     RL: DEV (Device component use); MOA (Modifier or additive use); RCT
     (Reactant); RACT (Reactant or reagent); USES (Uses)
        (in immobilization of hybridization of probes on MALDI
       matrixes; method for identifying nucleic acids by means of
       matrix-assisted laser desorption/ionisation mass spectrometry using
        charge tag-labeled hybridization probes)
     1519-55-7
RN
RN 227080-33-3
RN 227080-34-4
RN 64-17-5
RN
   67-56-1
RN
    67-63-0
RN
     67-64-1
    75-05-8
RN
    7732-18-5
RN
RN
     58-85-5
   9013-20-1
RN
L13 ANSWER 8 OF 15 CAPLUS COPYRIGHT 2003 ACS
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2001:936967 CAPLUS

AN

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DN
     136:309566
    A hydrophobic gel for epoxidation of olefins with organic peroxides
TТ
     Karli, Alfonsus; Larsen, Gustavo
ΑU
     Department of Chemical Engineering, University of Nebraska, Lincoln, NE,
CS
     68588-0126, USA
     Catalysis Letters (2001), 77(1-3), 107-111
SO
     CODEN: CALEER; ISSN: 1011-372X
    Kluwer Academic/Plenum Publishers
PΒ
DT
    Journal
LA
    English
    22-7 (Physical Organic Chemistry)
CC
     Section cross-reference(s): 21, 67
     The epoxidn. of cyclooctene by tert-Bu hydroperoxide (t-BuOOH) over a
AΒ
     TiO2-SiO2 xerogel made from a permethylated cyclooligosiloxane,
     tetraethylorthosilicate, and a Ti chloroalkoxide, was carried out in
     acetonitrile as the reaction solvent. The org. moieties of the
     hybrid gel (in this case, -Me groups) and the Ti content of the
     catalyst appear to be stable on prolonged exposure to the reaction medium.
     Besides very good stability, the hybrid catalyst in this study
     displays 100% selectivity toward cyclooctene epoxide prodn. Solid stafte
     29Si magic angle spinning NMR (29Si MASNMR), diffuse reflectance FTIR
     spectroscopy (DRIFTS), and chem. anal. were used to monitor the stability
     of the hybrid material. The kinetics of olefin epoxidn. was
     studied in a batch reactor in the 313-343 K range.
     hydrophobic epoxidn olefin org peroxide
ST
    MAS NMR spectroscopy
TΤ
        (CP, hydrophobic gel 29Si; hydrophobic gel for epoxidn. of olefins with
        org. peroxides)
     Diffuse reflectance IR spectroscopy
TΤ
        (Fourier-transform, hydrophobic gel; hydrophobic gel for epoxidn. of
        olefins with org. peroxides)
     Polysiloxanes, reactions
IΤ
     RL: CAT (Catalyst use); RCT (Reactant); RACT (Reactant or reagent); USES
     (Uses)
        (gel precursor; hydrophobic gel for epoxidn. of olefins with org.
        peroxides)
ΙT
     Activation energy
     Epoxidation
     Epoxidation catalysts
     Epoxidation kinetics
     Transition state structure
        (hydrophobic gel for epoxidn. of olefins with org. peroxides)
ΙT
     Alkenes, reactions
     Hydroperoxides
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (hydrophobic gel for epoxidn. of olefins with org. peroxides)
IT
     MAS NMR spectroscopy
        (hydrophobic gel 29Si; hydrophobic gel for epoxidn. of olefins with
        org. peroxides)
IT
     IR spectra
        (hydrophobic gel; hydrophobic gel for epoxidn. of olefins with org.
        peroxides)
TТ
     Gels
        (hydrophobic; hydrophobic gel for epoxidn. of olefins with org.
        peroxides)
     Peroxides, reactions
TΤ
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (org.; hydrophobic gel for epoxidn. of olefins with org. peroxides)
ΙT
     Adsorption
        (oxidant on gel; hydrophobic gel for epoxidn. of olefins with org.
        peroxides)
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TΤ
    Methyl group
        (surface tied to hybrid hydrophobic; hydrophobic gel for
        epoxidn. of olefins with org. peroxides)
     75-05-8, Acetonitrile, uses
TΤ
     RL: NUU (Other use, unclassified); USES (Uses)
        (better solvent than alcs.; hydrophobic gel for epoxidn. of olefins
        with org. peroxides)
     67-63-0, Isopropanol, reactions 7550-45-0, Titanium
ΙT
     tetrachloride, reactions
     RL: CAT (Catalyst use); RCT (Reactant); RACT (Reactant or reagent); USES
        (conversion to hydrophobic gel precursor; hydrophobic gel for epoxidn.
        of olefins with org. peroxides)
IT
     7722-84-1, Hydrogen peroxide, reactions
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (generation of oxygen in side reaction; hydrophobic gel for epoxidn. of
        olefins with org. peroxides)
     75-91-2, tert-Butyl hydroperoxide
                                        931-87-3, cis-Cyclooctene
TΤ
     RL: CPS (Chemical process); PEP (Physical, engineering or chemical
     process); PRP (Properties); RCT (Reactant); PROC (Process); RACT (Reactant
     or reagent)
        (hydrophobic gel for epoxidn. of olefins with org. peroxides)
     286-62-4, Cyclooctene epoxide
ΙT
     RL: FMU (Formation, unclassified); FORM (Formation, nonpreparative)
        (hydrophobic gel for epoxidn. of olefins with org. peroxides)
                   541-05-9
ΙT
     78-10-4, TEOS
     RL: CAT (Catalyst use); RCT (Reactant); RACT (Reactant or reagent); USES
        (hydrophobic gel precursor; hydrophobic gel for epoxidn. of olefins
        with org. peroxides)
                               75-65-0, tert-Butanol, uses
ΙT
     67-56-1, Methanol, uses
     RL: NUU (Other use, unclassified); USES (Uses)
        (oxidn. of solvent; hydrophobic gel for epoxidn. of olefins with org.
              THERE ARE 21 CITED REFERENCES AVAILABLE FOR THIS RECORD
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(21) Vu, D; Thesis University of Nebraska-Lincoln 1999
    75-05-8
RN
RN
     67-63-0
     7550-45-0
RN
     7722-84-1
RN
    75-91-2
RN
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286-62-4
RN
    78-10-4
RN
    541-05-9
RN
    67-56-1
RN
    75-65-0
RN
L13 ANSWER 9 OF 15 CAPLUS COPYRIGHT 2003 ACS
    2000:517764 CAPLUS
DN
    133:274951
    Combinatorial methods in sol-gel technology
TΙ
    Rantala, Juha T.; Kololuoma, Terho; Kivimaki, L.
ΑU
    VTT Electronics/Infotech Oulu, Oulu, Finland
CS
    Proceedings of SPIE-The International Society for Optical Engineering
SO
     (2000), 3941 (Combinatorial and Composition Spread Techniques in Materials
     and Device Development), 11-18
     CODEN: PSISDG; ISSN: 0277-786X
    SPIE-The International Society for Optical Engineering
PB
DT
    Journal
    English
LA
CC
    76-5 (Electric Phenomena)
     Section cross-reference(s): 56
     Sol-gel processing consists several variable parameters during materials
AΒ
     synthesis and post processing steps. The sol-gel synthesis is rather
     sensitive for the parameters such as pH, temp., type of catalyst, reaction
     time etc. However, this sensitivity can be taken as an advantage when
     developing and studying new materials and their properties. Furthermore,
     since the sol-gel technol. mainly describes the fabrication of solid state
     materials from a liq. phase by applying metal alkoxides or metal salts as
     precursors, the post processing such as sintering has crit. effects on the
     final form and properties of the solid material. Combinatorial chem. and
     methods are valuable tools to est. the effects of different variables and
     to build-up combinatorial libraries for the sol-gel technique. This paper
     generally describes potentials and the usage motivation of combinatorial
     chem. in the sol-gel technol. by taking into account some major steps in
     the synthesis and processing which are valuable for the estn. of the final
     product properties. Different kind of post processing steps in the
     combinatorial manner are studied in details. As an example the post
     processing of sol-gel derived semiconductor oxides and photosensitivity of
     hybrid sol-gel glasses are presented. The combinatorial treatment
     and measurement methods for these materials are explained.
     photocond antimony doped sol gel tin oxide glass; metal alkoxide salt
     sintering semiconductor oxide
     Semiconductor films
TΤ
        (Sb doped SnO2; photocond. of)
IT
     Dopants
        (Sb; effects on photocond. of Sb doped sol-gel SnO2)
ΙT
     Detergents
        (Triton X100; in sol-gel processing of SnO2)
     Refractive index
ΙT
        (and photocond. of Sb doped sol-gel SnO2)
ΙT
     Coating process
        (dip; in sol-gel processing of SnO2)
ΙT
        (films; effects on photocond. of Sb doped sol-gel SnO2)
ΙT
     Combinatorial chemistry
     Hydrolysis
     Photolithography
     Sintering
        (in sol-gel processing of SnO2)
     Alkali metal salts
ΙT
     Metal alkoxides
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ŔŊ

931-87-3

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RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical
     process); PROC (Process); USES (Uses)
        (in sol-gel processing of SnO2)
ΙT
     Glass substrates
        (lime; in photocond. of Sb doped sol-gel SnO2)
IT
     Composites
        (mol.; photocond. of Sb doped sol-gel SnO2)
     Photoconductivity
ΙT
     Sol-gel processing
        (of Sb doped sol-gel SnO2)
IT
     Crosslinking
        (photochem.; in sol-gel processing of SnO2)
     Electric conductivity
IΤ
        (photocond. of Sb doped sol-gel SnO2)
IT
     Zirconates
     Zirconates
     RL: PEP (Physical, engineering or chemical process); PRP (Properties); SPN
     (Synthetic preparation); PREP (Preparation); PROC (Process)
        (siloxane-, organically modified; photocond. of Sb doped
        sol-gel SnO2)
ÍΤ
     Coating process
        (spin; in sol-gel processing of SnO2)
     Coating process
ΤT
        (spray; in sol-gel processing of SnO2)
     Borosilicate glasses
TΤ
     RL: PEP (Physical, engineering or chemical process); TEM (Technical or
     engineered material use); PROC (Process); USES (Uses)
        (substrate; in photocond. of Sb doped sol-gel SnO2)
IT
     Polysiloxanes, properties
       Polysiloxanes, properties
     RL: PEP (Physical, engineering or chemical process); PRP (Properties); SPN
     (Synthetic preparation); PREP (Preparation); PROC (Process)
        (zirconate-, organically modified; photocond. of Sb doped sol-gel SnO2)
ΤТ
     7732-18-5, Water, processes
     RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical
     process); PROC (Process); USES (Uses)
        (deionized; in sol-gel processing of SnO2)
     9002-93-1, Triton X100
IΤ
     RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical
     process); PROC (Process); USES (Uses)
        (detergent; in sol-gel processing of SnO2)
ΙT
     10025-91-9, Antimony chloride (SbCl3)
     RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical
     process); PROC (Process); USES (Uses)
        (dopant source; in sol-gel processing of SnO2)
     7440-36-0, Antimony, uses
TΤ
     RL: MOA (Modifier or additive use); USES (Uses)
        (dopant; effects on photocond. of Sb doped sol-gel SnO2)
     67-63-0, Isopropanol, processes 67-64-1, Acetone, processes
ΙT
     79-41-4, Methacrylic acid, processes
                                            2171-98-4, Zirconium isopropoxide
                 24623-80-1, Tin chloride (SnCl4) tetrahydrate
     2530-85-0
     RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical
     process); PROC (Process); USES (Uses)
        (in sol-gel processing of SnO2)
     162881-26-7, Irgacure 819
IT
     RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical
     process); PROC (Process); USES (Uses)
        (photo initiator; in sol-gel processing of SnO2)
ΙT
     7440-37-1, Argon, processes
     RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical
     process); PROC (Process); USES (Uses)
        (plasma treatment; effects on photocond. of Sb doped sol-gel SnO2)
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ΙT
   1332-29-2P, Tin oxide
     RL: PEP (Physical, engineering or chemical process); PRP (Properties); SPN
     (Synthetic preparation); PREP (Preparation); PROC (Process)
        (sol-gel; photocond. of)
              THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE
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(2) Kololuoma, T; SPIE Photonic West 1999, Pl34 CAPLUS
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   7732-18-5
RN
   9002-93-1
RN 10025-91-9
RN 7440-36-0
RN
   67-63-0
   67-64-1
RN
    79-41-4
RN
   2171-98-4
RN
   2530-85-0
RN
   24623-80-1
RN
    162881-26-7
RN
RN
    7440-37-1
RN
   1332-29-2P
L13 ANSWER 10 OF 15 CAPLUS COPYRIGHT 2003 ACS
    1999:206999 CAPLUS
DN
    130:268425
    Rubber-ceramic composites and their manufacture
TТ
   Hamanaka, Seiko; Nishimoto, Kazuo
IN
   Nichias Corp., Japan
PA
SO
     Jpn. Kokai Tokkyo Koho, 7 pp.
     CODEN: JKXXAF
DT
    Patent
LA
     Japanese
    ICM C08J003-20
IC
     ICS C08K003-22; C08L021-00; C08L083-04; C08C019-26
     39-9 (Synthetic Elastomers and Natural Rubber)
     Section cross-reference(s): 57
FAN.CNT 1
                                          APPLICATION NO. DATE
                  KIND DATE
     PATENT NO.
                     ____
                                           _____

      JP 11080373
      A2
      19990326

      JP 3295023
      B2
      20020624

                                          JP 1997-242752 19970908
PΤ
PRAI JP 1997-242752
                           19970908
     Rubbers dissolved in org. solvents or liq. rubbers are mixed with
     organometallic compds. and then subjected to sol-gel reaction so that
     oxides of the metals derived from the organometallic compds. are formed
     and dispersed finely, polymn. occurs, and chem. bonds are formed between
     the organometallic compds. and rubbers. Thus, SBR rubber was dissolved in
     THF, mixed with 120.0 phr (.gamma.-isocyanatopropyl)triethoxysilane,
     gelled with H2O, ammonia, and NH4F, dried, further mixed with S 1.5, ZnO
     5.0, and vulcanizing accelerators 3.0 phr, and press-vulcanized to give a
     sheet showing improved heat resistance compared to an SBR sheet contg.
     19.0 phr white carbon.
     SBR rubber ceramic composite manuf; isocyanatopropyltriethoxysilane rubber
ST
     composite heat resistance
     Cold-resistant materials
ΤТ
     Heat-resistant materials
       Hybrid organic-inorganic materials
     Sol-gel processing
     Transparent materials
        (manuf. of rubber-ceramic composites by sol-gel process)
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TΥ
    Oxides (inorganic), preparation
    RL: IMF (Industrial manufacture); PRP (Properties); PREP (Preparation)
       (manuf. of rubber-ceramic composites by sol-gel process)
    EPDM rubber
ΤТ
    Silicone rubber, preparation
    Styrene-butadiene rubber, preparation
    RL: IMF (Industrial manufacture); PRP (Properties); PREP (Preparation)
       (reaction products, with organometallic compds.; manuf. of
       rubber-ceramic composites by sol-gel process)
    Polysiloxanes, preparation
ΙΤ
    RL: IMF (Industrial manufacture); PRP (Properties); PREP (Preparation)
       (reaction products, with rubber; manuf. of rubber-ceramic composites by
       sol-gel process)
    67-63-ODP, Isopropanol, polytitanate derivs., reaction products
TΤ
                  546-68-9DP, Titanium tetraisopropoxide, reaction products
    with rubber 2269-22-9DP, reaction products with rubber 4420-74-0DP,
     (.gamma.-Mercaptopropyl)trimethoxysilane, reaction products with rubber
    9016-00-6DP, Dimethyl siloxane, sru, reaction products with
            24801-88-5DP, reaction products with rubber
                                                         31900-57-9DP,
     Dimethylsilanediol homopolymer, reaction products with rubber
     40372-72-3DP, Bis(3-triethoxysilylpropyl)tetrasulfane, reaction products
    with rubber
    RL: IMF (Industrial manufacture); PRP (Properties); PREP (Preparation)
       (manuf. of rubber-ceramic composites by sol-gel process)
IT
    9003-55-8P
    RL: IMF (Industrial manufacture); PRP (Properties); PREP (Preparation)
       (styrene-butadiene rubber, reaction products, with organometallic
       compds.; manuf. of rubber-ceramic composites by sol-gel process)
    67-63-0DP
RN
   546-68-9DP
RN
   2269-22-9DP
RN
RN
   4420-74-0DP
RN 9016-00-6DP
   24801-88-5DP
   31900-57-9DP
RN 40372-72-3DP
RN 9003-55-8P
L13 ANSWER 11 OF 15 CAPLUS COPYRIGHT 2003 ACS
   1999:502748 CAPLUS
AN
    131:149863
DN
    Molecular sieving silica membrane fabrication process
ΤI
   Raman, Narayan K.; Brinker, Charles Jeffrey
TN
   Gas Research Institute, USA; Sandia National Laboratories
PA
SO
   U.S., 18 pp.
    CODEN: USXXAM
DT
    Patent
LA English
    ICM B05D005-00
TC
NCL 427244000
    66-4 (Surface Chemistry and Colloids)
     Section cross-reference(s): 36, 78
FAN.CNT 2
                    KIND DATE
                                         APPLICATION NO.
    PATENT NO.
                                         _____
     ______
                                       US 1998-13346∕
                                                          19980126
                    A 19990810
    US 5935646
PΙ
                    A 19980623
                                         US 1996-7027/45
                                                          19960823
    US 5770275
                          19960823
PRAI US 1996-702745
    A process for producing a mol. sieve silica membrane comprising depositing
     a hybrid org.-inorg. polymer comprising at least one org.
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constituent and at least one inorg. constituent on a porous substrate material and removing at least a portion of the at least one org.

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constituent of the hybrid org.-inorg. polymer, forming a porous
     film.
     silica membrane mol sieve org inorg polymer
ST
ΙT
     Permeation
        (permeance and pore size and structure properties of silica membranes
        prepd. by copolymn. of TEOS and MTES and deposited on a porous alumina
    Polysiloxanes, properties
IT
    RL: PEP (Physical, engineering or chemical process); PRP (Properties); SPN
     (Synthetic preparation); PREP (Preparation); PROC (Process)
        (permeance and pore size and structure properties of silica membranes
        prepd. by copolymn. of TEOS and MTES and deposited on a porous alumina
        substrate)
    Molecular sieves
ΤТ
     Pore size
     Pore size distribution
     Pore structure
     Porosity
     Porous materials
        (permeance and pore size and structure properties of silica membranes
        prepd. by copolymn. of TEOS and MTES detd. by mol. sieving of various
ΙT
    Xerogels
        (permeance and pore size and structure properties of silica membranes
        prepd. by copolymn. of TEOS and MTES xerogels)
     67-63-0, Isopropanol, uses
                                 75-65-0, tert-Butanol, uses
TT
     27129-87-9, 3,5-Dimethyl benzyl alcohol
     RL: NUU (Other use, unclassified); USES (Uses)
        (mol. probing of surface derivatized silica membranes of TEOS and MTES
        with)
     1344-28-1, Alumina, uses
IT
     RL: NUU (Other use, unclassified); USES (Uses)
        (permeance and pore size and structure properties of silica membranes
        prepd. by copolymn. of TEOS and MTES and deposited on a porous alumina
        substrate)
                                 2031-67-6, Methyltriethoxysilane
     78-10-4, Tetraethoxysilane
TΤ
     RL: PEP (Physical, engineering or chemical process); PRP (Properties); RCT
     (Reactant); PROC (Process); RACT (Reactant or reagent)
        (permeance and pore size and structure properties of silica membranes
        prepd. by copolymn. of TEOS and MTES and deposited on a porous alumina
        substrate)
     88029-70-3P
ΙT
     RL: PEP (Physical, engineering or chemical process); PRP (Properties); SPN
     (Synthetic preparation); PREP (Preparation); PROC (Process)
        {permeance and pore size and structure properties of silica membranes
        prepd. by copolymn. of TEOS and MTES and deposited on a porous alumina
        substrate)
     74-82-8, Methane, properties
                                    124-38-9, Carbon dioxide, properties
ΤT
                                      7440-59-7, Helium, properties
     2551-62-4, Sulfur hexafluoride
     7727-37-9, Nitrogen, properties
     RL: PEP (Physical, engineering or chemical process); PRP (Properties);
     PROC (Process)
        (permeance and pore size and structure properties of silica membranes
        prepd. by copolymn. of TEOS and MTES detd. by mol. sieving of)
     2996-92-1, Phenyltrimethoxysilane
ΙT
     RL: MOA (Modifier or additive use); PEP (Physical, engineering or chemical
     process); PRP (Properties); RCT (Reactant); PROC (Process); RACT (Reactant
     or reagent); USES (Uses)
        (surface derivatization of silica membranes prepd. by copolymn. of TEOS
        and MTES detd. by addn. of during the polymn. process)
             THERE ARE 15 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT
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RF.

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    V125, P1
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(13) Pasco; US 4221748 1980
(14) Raman; US 5770275 1998 CAPLUS
(15) Robeson, L; Journal of Membrane Science 1991, V62, P165 CAPLUS
RN
     67-63-0
RN
     75-65-0
RN
   27129-87-9
RN
    1344-28-1
    78-10-4
RN
    2031-67-6
RN
    88029-70-3P
RN
    74-82-8
RN
    124-38-9
RN
    2551-62-4
RN
     7440-59-7
RN
     7727-37-9
RN
     2996-92-1
RN
L13 ANSWER 12 OF 15 CAPLUS COPYRIGHT 2003 ACS
    1998:66185 CAPLUS
ΑN
DN
    128:116668
    Composite pervaporation membrane with ceramic support structure
TΙ
PΑ
    Mauz, Matthias, Germany
SO
     Ger. Offen., 10 pp.
     CODEN: GWXXBX
DT
     Patent
     German
LA
     ICM B01D069-12
TC
     ICS B01D061-36; B01D001-00; B01D005-00; C07B063-00
     47-2 (Apparatus and Plant Equipment)
     Section cross-reference(s): 38, 57, 80
FAN.CNT 1
                      KIND DATE
     PATENT NO.
                                           APPLICATIÓN NO.
                      ____
                                           DE 1996-19629061 19960719
     DE 19629061
                      Αl
                            19980122
                            19960719
PRAI DE 1996-19629061
     The composite membrane is a pore-free polymer cover layer on a porous
     ceramic support layer having pore size 0.005-0.5 .mu.m. The org.
     solvent-selective polymer cover layer can be a
     polydimethylsiloxane, polyamide, or polyether. The sides of the
     ceramic layer are sealed with an epoxy resin or polyurethane. The
     membrane has good thermal and chem. stability, and can be used for
     pervaporation of solvent mixts. in an anal. lab., e.g., HPLC solvents. It
     may be used for hybrid distn.-thermal sepn. methods.
    pervaporation membrane ceramic support analytical solvent
ST
ΙT
     Ceramic membranes
     Solvents
        (composite pervaporation membrane with ceramic support structure)
ΙT
     Polyamides, uses
     Polyethers, uses
```

```
(composite pervaporation membrane with ceramic support structure)
      Epoxy resins, uses
 IΤ
      RL: NUU (Other use, unclassified); USES (Uses)
         (composite pervaporation membrane with ceramic support structure)
 ΙT
      Polyurethanes, uses
      RL: NUU (Other use, unclassified); USES (Uses)
         (composite pervaporation membrane with ceramic support structure)
 ΙT
      Polysiloxanes, uses
      RL: DEV (Device component use); USES (Uses)
         (di-Me; composite pervaporation membrane with ceramic support
         structure)
 TΤ
      Pervaporation
         (membranes; composite pervaporation membrane with ceramic support
 IT
      56-81-5, Glycerin, properties 64-18-6, Formic acid, properties
      67-63-0, Isopropanol, properties 67-64-1, Acetone, properties 77-92-9, Citric acid, properties 141-78-6, Ethyl acetate, properties
      7647-01-0, Hydrochloric acid, properties 7664-93-9, Sulfuric acid,
      properties
      RL: PRP (Properties)
         (composite pervaporation membrane with ceramic support structure)
 IΤ
      64-17-5P, Ethanol, preparation 64-19-7P, Acetic acid, preparation
      67-56-1P, Methanol, preparation 75-05-8P, Acetonitrile, preparation
      142-82-5P, Heptane, preparation 7732-18-5P, Water, preparation
      RL: PUR (Purification or recovery); PREP (Preparation)
         (composite pervaporation membrane with ceramic support structure)
 RN
      56-81-5
 RN
      64-18-6
 RN
     67-63-0
RN
     67-64-1
RN
     77-92-9
RN
     141-78-6
RN
     7647-01-0
RN
     7664-93-9
RN
    64-17-5P
RN
    64-19-7P
RN
     67-56-1P
RN
     75-05-8P
RN
     142-82-5P
RN
     7732-18-5P
L13 ANSWER 13 OF 15 CAPLUS COPYRIGHT 2003 ACS
AN
     1998:41700 CAPLUS
DN
     128:119648
TI
     Extraction process for producing poly(DL-lactide-glycolide) microspheres
     Van Hamont, John; Thies, Curt; Reid, Robert H.; McQueen, Charles E.;
IN
     Setterstrom, Jean A.
PΑ
     United States Dept. of the Army, USA
     U.S., 9 pp., Cont.-in-part of U.S. Ser. No. 242,960.
SO
     CODEN: USXXAM
DT
     Patent
LA
    English
TC
    ICM A61K009-50
NCL 424501000
CC
     63-6 (Pharmaceuticals)
FAN.CNT 13
    PATENT NO.
                     KIND DATE
                                           APPLICATION NO. DATE
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                           -----
                                           -----
    US 5705197
                     А
                           19980106
                                          US 1996-698896 19960816
                     A
B1
    US 5693343
                           19971202
                                          US 1994-242960 19940516
    US 6447796
                           20020910
                                         US 1997-920326 19970821
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RL: DEV (Device component use); USES (Uses)

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PRAI US 1994-242960
                      A2
                           19940516
                           19840316
    US 1984-590308
                      A2
                           19900315
                     B2
    US 1990-493597
                     В2
                           19900511
    US 1990-521945
                     B2
                           19910424
    US 1991-690485
                     B2 19911121
    US 1991-805721
                     A2 19920410
    US 1992-867301
                     A2 19950522
    US 1995-446148
    US 1995-446149 B2 19950522
    US 1996-590973 B2 19960124
                    A2 19960705
    US 1996-675895
    US 1996-698896 A2 19960816
                     A2
                          19970127
    US 1997-789734
    A hybrid evapn.-extn. process for prepg. microspheres of a
AΒ
     poly(DL-lactide-glycolide) (I) biodegradable polymer, comprising: (a)
     prepg. a lyophilized biol. active material-sucrose matrix; adding
     acetonitrile solvent to biol. active material-sucrose matrix to form a
     soln.; (b) prepg. a soln. of a biodegradable I by adding acetonitrile
     solvent to the polymer; (c) adding the biodegradable I acetonitrile soln.
     to the biol. active material-sucrose acetonitrile soln.; (d) adding with
     stirring an oil contg. lecithin to the I-sucrose-biol. active material
     soln. to evap. acetonitrile and form an emulsion contg. microspheres of I
     biodegradable polymers; (e) adding the emulsion from step d. into a
     solvent selected from heptane, hexane, pentane or isopropanol; and (f)
     collecting microspheres of I biodegradable polymers of from 1.0 to about
     10.0 .mu.m after filtration and washing with a fresh solvent selected from
     heptane, hexane, pentane or isopropanol. I microspheres between
     5-10.mu.m, with very little aggregation were prepd. according to above
     pharmaceutical microsphere extn polylactide polyglycolide
ST
IΤ
    Antigens
     Lecithins
     Paraffin oils
       Polysiloxanes, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (extn. process for producing poly(DL-lactide-glycolide) microspheres)
ΙT
     Drug delivery systems
        (freeze-dried; extn. process for producing poly(DL-lactide-glycolide)
        microspheres)
IT
     Lubricating oils
        (machine; extn. process for producing poly(DL-lactide-glycolide)
        microspheres)
IT
     Drug delivery systems
        (microspheres; extn. process for producing poly(DL-lactide-glycolide)
        microspheres)
     57-50-1, biological studies 67-63-0, Isopropanol;, biological
ΙT
               75-05-8, Acetonitrile, biological studies 109-66-0, Pentane,
     studies
                         110-54-3, Hexane, biological studies
                                                               142-82-5,
     biological studies
                                   26780-50-7, Poly(DL-lactide-glycolide)
     Heptane, biological studies
     RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
        (extn. process for producing poly(DL-lactide-glycolide) microspheres)
RN
     57-50-1
     67-63-0
RN
     75-05-8
RN
     109-66-0
RN
RN
     110-54-3
     142-82-5
RN
     26780-50-7
RN
L13 ANSWER 14 OF 15 CAPLUS COPYRIGHT 2003 ACS
NΑ
     1998:539347 CAPLUS
DN
     129:277633
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Meerwein-Ponndorf-Verley reductions mediated by lanthanide-alkoxide-
TI
    functionalized mesoporous silicates
    Anwander, R.; Palm, C.
ΑU
    Institut fur Technische Chemie I, Universitat Stuttgart, Stuttgart,
CS
    D-70569, Germany
    Studies in Surface Science and Catalysis (1998), 117 (Mesoporous Molecular
SO
    Sieves 1998), 413-420
    CODEN: SSCTDM; ISSN: 0167-2991
    Elsevier Science B.V.
PΒ
DT
    Journal
    English
LA
    45-4 (Industrial Organic Chemicals, Leather, Fats, and Waxes)
    Section cross-reference(s): 24, 67
    Lanthanide alkoxide moieties were grafted onto mesoporous silicate MCM-41
     via surface siloxide linkage. Two immobilization procedures
     were used: treatment of dehydrated MCM-41 with rare earth alkoxide solns.
     and prepn. of silylamide derivs. in heterogeneous media. The latter
     procedure facilitated introduction of a wide range of alkoxide ligands via
     ligand exchange reactions of MCM-41/rare earth amide hybrid
     species with various alcs. The materials were characterized by elemental
     anal. and FTIR spectroscopy. Nitrogen physisorption measurements revealed
     that the effective mean pore diam. and pore vol. were markedly affected by
     the steric bulk of alkoxide ligands. The hybrid materials are
     efficient precatalysts in the Meerwein-Ponndorf-Verley redn. of
     4-tert-butylcyclohexanone to the corresponding 4-tert-butylcyclohexanol
     isomers. The transformations were conducted in 2-propanol and conversions
     >95 - were accomplished within five hours even at ambient temp., and
     without formation of any side products.
     lanthanide alkoxide zeolite catalyst redn; ketone redn alc lanthanide
ST
     mesoporous silicate; Meerwein Ponndorf Verley redn silicate catalyst
ΙT
     Rare earth complexes
     RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation);
     USES (Uses)
        (alc.; neodymium alkoxide-functionalized mesoporous zeolite catalysts
        for Meerwein-Ponndorf-Verley redn. ketones to alcs.)
     Zeolite MCM-41
ΙT
     RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation);
        (lanthanide-alkoxide-functionalized; neodymium alkoxide-functionalized
        mesoporous zeolite catalysts for Meerwein-Ponndorf-Verley redn. ketones
        to alcs.)
IT
     Porous materials
        (mesoporous; neodymium alkoxide-functionalized mesoporous zeolite
        catalysts for Meerwein-Ponndorf-Verley redn. ketones to alcs.)
ΙT
     Dehydration reaction
     Physisorption
     Reduction
     Substitution reaction, coordinative
        (neodymium alkoxide-functionalized mesoporous zeolite catalysts for
        Meerwein-Ponndorf-Verley redn. ketones to alcs.)
     Alcohols, preparation
TT
     RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation);
     USES (Uses)
        (rare earth complexes; neodymium alkoxide-functionalized mesoporous
        zeolite catalysts for Meerwein-Ponndorf-Verley redn. ketones to alcs.)
     41836-23-1DP, Tris[bis(trimethylsilyl)amido]neodymium, mesoporous silicate
               142042-03-3DP, mesoporous silicate derivs.
     RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation);
     USES (Uses)
        (neodymium alkoxide-functionalized mesoporous zeolite catalysts for
        Meerwein-Ponndorf-Verley redn. ketones to alcs.)
     98-52-2P, 4-tert-Butylcyclohexanol
TΤ
```

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RL: IMF (Industrial manufacture); PREP (Preparation)
        (neodymium alkoxide-functionalized mesoporous zeolite catalysts for
        Meerwein-Ponndorf-Verley redn. ketones to alcs.)
    64-17-5, Ethanol, reactions 67-56-1, Methanol, reactions 67-63-0
TΤ
     , Isopropyl alcohol, reactions
                                      75-84-3, Neopentyl alcohol
     4-tert-Butylcyclohexanone
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (neodymium alkoxide-functionalized mesoporous zeolite catalysts for
        Meerwein-Ponndorf-Verley redn. ketones to alcs.;
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     98-53-3
L13 ANSWER 15 OF 15 CAPLUS COPYRIGHT 2003 ACS
     1999:208880 CAPLUS
AN
DN
     130:313504
     Fabric softening and antistatic agents containing N-
ΤI
     alkanolalkylenepolyamine ester amide compounds
     Inoue, Kimi
IN
     Kao Corp., Japan
PΑ
     Jpn. Kokai Tokkyo Koho, 12 pp.
SO
     CODEN: JKXXAF
DT
     Patent
LA
     Japanese
     ICM D06M013-46
TC
     46-5 (Surface Active Agents and Detergents)
     Section cross-reference(s): 40
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FAN.CNT 1

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APPLICATION NO. DATE
     PATENT NO. KIND DATE
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                                           JP 1997-235229 19970829

      JP 11081134
      A2 19990326

      JP 3346235
      B2 20021118

PΙ
JP 3346235 B2 20021118
PRAI JP 1997-235229 19970829
    MARPAT 130:313504
    The agents comprise (A) R1N(CmH2mOCOR2)(CnH2nNHCOR3) (R1 = C1-4 alkyl,
     hydroxyalkyl; R2, R3 = C11-21 alkyl or alkenyl; m = 1-10; n = 2-3), their
     neutralized products or quaternary ammonium compds.; (B) C12-22 linear or
     branched (un) satd. carboxylic acids; (C) C2-6 glycols, C3-6 aliph. alcs.,
     C8-18 arom. esters or/and C10-15 terpenoid compds.; and (D) perfume.
     Thus, an antistatic and softening agent was obtained from a mixt. of
     N-methyl-N-(hydrogenated tallow fatty acid esterified hydroxyethyl)-N-
     (hydrogenated tallow fatty acid amidated aminopropyl)amine.cntdot.HCl salt
     5, hydrogenated tallow fatty acid 1, a 50:25:10:15 mixt. of di-Et
     phthalate, benzyl salicylate, benzyl acetate and citronellyl acetate, 0.1,
     and a perfume 0.03.
     fabric antistatic softening agent quaternary ammonium compd; fatty acid
ST
     antistatic softening fabric; perfume antistatic softening fabric;
     hydrogenated tallow fatty acid alkanolamide softening fabric
ΙT
     Alcohols, uses
     RL: TEM (Technical or engineered material use); USES (Uses)
        (aliph.; fabric softening and antistatic agents from ammonium compds.)
ΤТ
     Esters, uses
     RL: TEM (Technical or engineered material use); USES (Uses)
        (arom.; fabric softening and antistatic agents from ammonium compds.)
     Quaternary ammonium compounds, uses
IΤ
     Terpenes, uses
     RL: TEM (Technical or engineered material use); USES (Uses)
        (fabric softening and antistatic agents contg. N-
        alkanolalkylenepolyamine ester amide compds.)
     Antistatic agents
TΤ
     Fabric softeners
     Perfumes
        (fabric softening and antistatic agents from ammonium compds.)
     Polysiloxanes, uses
ΙT
     RL: MOA (Modifier or additive use); USES (Uses)
        (fabric softening and antistatic agents from ammonium compds.)
     Carboxylic acids, uses
IT
     RL: TEM (Technical or engineered material use); USES (Uses)
        (fabric softening and antistatic agents from ammonium compds.)
TT
     Essential oils
     RL: TEM (Technical or engineered material use); USES (Uses)
        (lavender, perfume; fabric softening and antistatic agents from
        ammonium compds.)
     Lavender (Lavandula hybrida)
ΙT
        (oils, perfume; fabric softening and antistatic agents from ammonium
        compds.)
     Essential oils
IΤ
     RL: TEM (Technical or engineered material use); USES (Uses)
        (orange, sweet, perfume compn.; fabric softening and antistatic agents
        contg. N-alkanolalkylenepolyamine ester amide compds.)
     Fatty acids, uses
ΙT
     RL: TEM (Technical or engineered material use); USES (Uses)
        (palm-oil, esters, compds. with N-alkyl-N-ethanol-1,3-propylenediamine,
        salts or quaternary compds.; fabric softening and antistatic agents
        contg. N-alkanolalkylenepolyamine ester amide compds.)
ΙT
     Fatty acids, uses
     RL: TEM (Technical or engineered material use); USES (Uses)
         'palm-oil; fabric softening and antistatic agents contg.
        N-alkanolalkylenepolyamine ester amide compds.)
ΙT
     Essential oils
```

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RL: TEM (Technical or engineered material use); USES (Uses)
        (perfume compn.; fabric softening and antistatic agents contg.
        N-alkanolalkylenepolyamine ester amide compds.)
TΤ
    Palm oil
     RL: TEM (Technical or engineered material use); USES (Uses)
        (stearins, compds. with N-alkyl-N-ethanol-1,3-propylenediamine, salts
        or quaternary compds.; fabric softening and antistatic agents contg.
        N-alkanolalkylenepolyamine ester amide compds.)
     Fatty acids, uses
ΙΤ
     RL: TEM (Technical or engineered material use); USES (Uses)
        (tallow, hydrogenated, esters, compds. with N-alkyl-N-ethanol-1,3-
        propylenediamine, salts or quaternary compds.; fabric softening and
        antistatic agents contg. N-alkanolalkylenepolyamine ester amide
        compds.)
     Fatty acids, uses
IT
     RL: TEM (Technical or engineered material use); USES (Uses)
        (tallow, hydrogenated; fabric softening and antistatic agents contg.
        N-alkanolalkylenepolyamine ester amide compds.)
     Fatty acids, uses
IT
     RL: TEM (Technical or engineered material use); USES (Uses)
        (tallow; fabric softening and antistatic agents contg.
        N-alkanolalkylenepolyamine ester amide compds.)
     41999-70-6D, compds. with fatty acids, salts or quaternary compds.
ΙT
     151955-40-7 161444-02-6 171064-63-4 171064-64-5 175716-84-4
     RL: TEM (Technical or engineered material use); USES (Uses)
        (fabric softening and antistatic agents contg. N-
        alkanolalkylenepolyamine ester amide compds.)
     112-85-6, Docosanoic acid 506-30-9, Eicosanoic acid 544-63-8,
ΙT
                               73756-39-5
     Tetradecanoic acid, uses
     RL: TEM (Technical or engineered material use); USES (Uses)
        (fabric softening and antistatic agents from ammonium compds.)
     67-56-1, Carbinol, uses 77-83-8, Aldehyde C16 78-69-3 78-70-6,
ΙT
     Linalool 79-77-6, .beta.-Ionone 80-54-6, Lilial 81-14-1, Musk ketone
     93-04-9, Yara yara 97-53-0, Eugenol 101-86-0, Hexyl cinnamic aldehyde 103-95-7, Cyclamen aldehyde 104-61-0, Aldehyde C18 106-02-5, Pentalide
                           106-24-1 110-41-8, Methylnonylacetaldehyde
     106-22-9, Citronellol
     120-57-0, Heliotropin 121-32-4, Ethylvanillin 121-33-5, Vanillin
                                  125-12-2, Isobornyl acetate
                                                                 127-48-0,
     123-11-5, Anisaldehyde, uses
            128-51-8, Nopyl acetate 151-05-3, Dimethylbenzylcarbinyl acetate
     Edion
                           1205-17-0, Helional
                                                 1506-02-1, Tentarome
     470-82-6, Eucalyptol
                                 5471-51-2, Raspberry ketone 6864-62-6,
     2050-08-0, Amyl salicylate
                          8000-41-7, Terpineol 16409-43-1, Rose oxide
     Phenyl acetoacetate
     23726-91-2, .beta.-Damascone
                                   27134-07-2
                                                 30385-25-2, Dihydromyrcenol
     32210-23-4, p-tert-Butylcyclohexyl acetate 32388-55-9, Acetylcedrene
                           55066-48-3, Phenoxanol
                                                     63429-28-7,
     41199-19-3, Ambrinol
                                                 80111-68-8, Damascone
                           68140-53-4, Fruitate
     .beta.-Methylionone
     80449-98-5, Liral 139504-68-0, Amber core 145334-39-0
                                                                176201-25-5,
                        176201-49-3, Poarenet
                                                  177771-82-3, Ambroxan
     Aldehyde C14 Peach
     183601-27-6
                  223447-73-2, Tetrahydromugol
     RL: TEM (Technical or engineered material use); USES (Uses)
        (perfume compn.; fabric softening and antistatic agents contg.
        N-alkanolalkylenepolyamine ester amide compds.)
     60-12-8, Phenylethyl alcohol 77-54-3, Cedryl acetate
                                                             80-26-2
ΙT
     91-64-5, Coumarin 93-08-3, Methyl .beta.-naphthyl ketone
     Diphenyl oxide 104-55-2, Cinnamic aldehyde 122-78-1, Phenyl
                   143-07-7, Dodecanoic acid, uses 497-62-1
                                                                 1222-05-5,
     acetaldehyde
                                            43052-87-5, .alpha.-Damascone
     Pearlide 21677-96-3, Geranylnitrile
                                                             124899-75-8
     51566-62-2, Citronellyl nitrile
                                       68039-49-6, Tripral
     188647-24-7
     RL: TEM (Technical or engineered material use); USES (Uses)
        (perfume compn.; fabric softening and antistatic agents from ammonium
        compds.)
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57-11-4, Octadecanoic acid, uses 57-55-6, 1,2-Propanediol, uses 84-66-2, Diethyl phthalate
         93-92-5, Styrallyl acetate 103-45-7 103-54-8, Cinnamyl acetate
        105-85-1, Citronellyl formate 105-87-3, Geranyl acetate 107-21-1, 1,2-Ethanediol, uses 112-80-1, Oleic acid, uses 115-95-7, Linalyl
        acetate 118-58-1, Benzyl salicylate 119-36-8, Methyl salicylate
        122-69-0, Cinnamyl cinnamate 134-20-3, Methyl anthranilate 140-11-4,
        Benzyl acetate 150-84-5, Citronellyl acetate
        acetate 928-96-1, cis-3-Hexenol 6259-76-3, Hexyl salicylate
                                                            326-61-4, Heliotropyl
        25265-71-8, Dipropylene glycol
                                           56539-66-3, 3-Methoxy-3-methylbutanol
        65405-77-8, cis-3-Hexenyl salicylate
        RL: TEM (Technical or engineered material use); USES (Uses)
           sperfume retention aids; fabric softening and antistatic agents contg.
           N-alkanolalkylenepolyamine ester amide compds.)
        57-10-3, Palmitic acid, uses
  ΙT
        RL: TEM (Technical or engineered material use); USES (Uses)
           (perfume retention aids; fabric softening and antistatic agents from
           ammonium compds.)
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